

Geochemical Perspectives Linking Arsenic Fate and Retention to Iron and Sulfur Cycling

Protect Your Health

The soils and sediments in this area contain harmful levels of lead and other metals. Small children and pregnant women are at the greatest risk from exposure.

KEEP CLEAN ! Wash your hands and face before you eat anything. Wash toys, bottles, and pacifiers if they have been in contact with soil or dust. Remove loose soil from your clothing, camping equipment, and pets before leaving the area. Wash all items when you return home.

EAT CLEAN ! Drink, cook, and wash only with water from home or other approved source. Do not use river water. Always eat at a table or clean surface off the ground. Clean fish thoroughly and eat only fish fillets.

PLAY CLEAN ! Children should play in grassy areas and avoid loose soil, dust, and muddy areas. No mud pies.

Healthy Choices.....Healthy Kids !

For more information call Panhandle Health District / Kellogg at (208) 783-0707

LEAD AND MINING

The Silver Valley is one of the oldest and largest mining communities in our country. The major metals that have been mined and smelted include silver, zinc and lead. Lead is a very common metal and is used extensively in many aspects of our daily lives including gasoline, batteries, solder, and paint. Lead has been around for over 400 years.

Our bodies do not need lead. Excessive absorption can result by either swallowing or breathing it. It is one of the most preventable childhood health problems of today.

For over 100 years these lower grounds have been contaminated by lead and other heavy metals which have been washed down by the river.

By following these simple guidelines you will keep contact to a minimum.

1. Wash your hands before eating.
2. Do not eat on the ground.
3. Do not play in the uncapped areas.
4. Do not drink the water in the river, even if filtered.

Please
**TAKE
ONE**

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Dartmouth College
Department of Earth Sciences

Funding:
NIEHS-SBRP
EPA
Environ Foundation

Arsenic in the Environment

- Arsenic not rare in the environment
 - “average soil”: about 10 mg As/kg
- Toxic environmental effects associated with arsenic not rare.
 - Effects of arsenic significant even at very low dissolved levels
 - Effects of arsenic are widespread

Cambodian rice field in As-impacted area

Natural Sources: Arsenic in Groundwater

- Arsenic concentrations in sediments in Bangladesh and Cambodia are not high. In fact, they are frequently **below** average.
- **Chemical Conditions** create elevated *dissolved* arsenic concentrations.
 - LANDFILLS (lined and unlined) are not unique, but are reactors in which pH and redox conditions are modulated by a combination of biological, chemical, and physical processes
 - **Microbes**
 - **Electron Source** (organic matter, H_2)
 - **Terminal Electron Acceptors** (Oxygen, Iron(III), Sulfate, CO_2)



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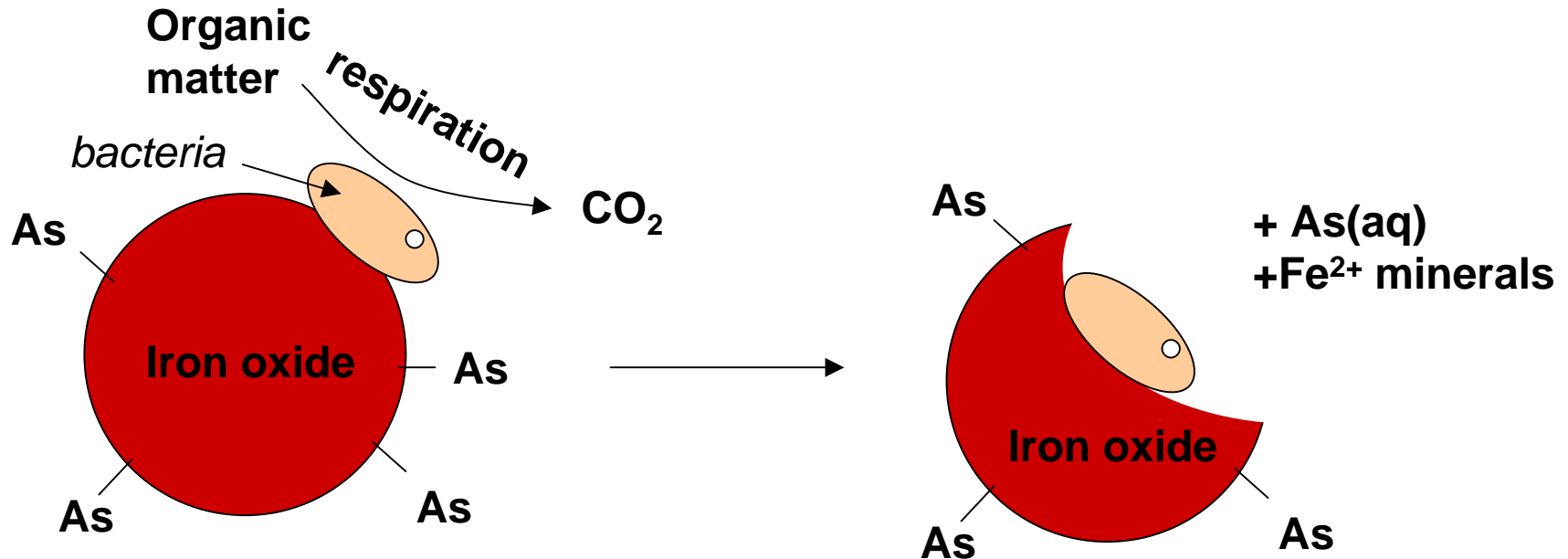
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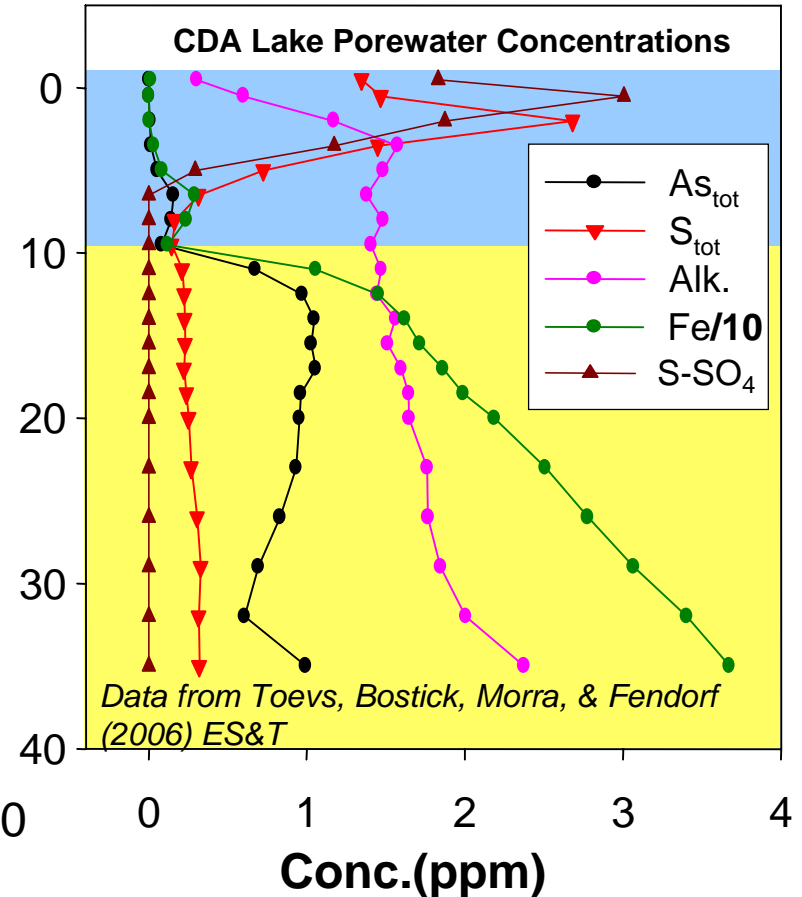
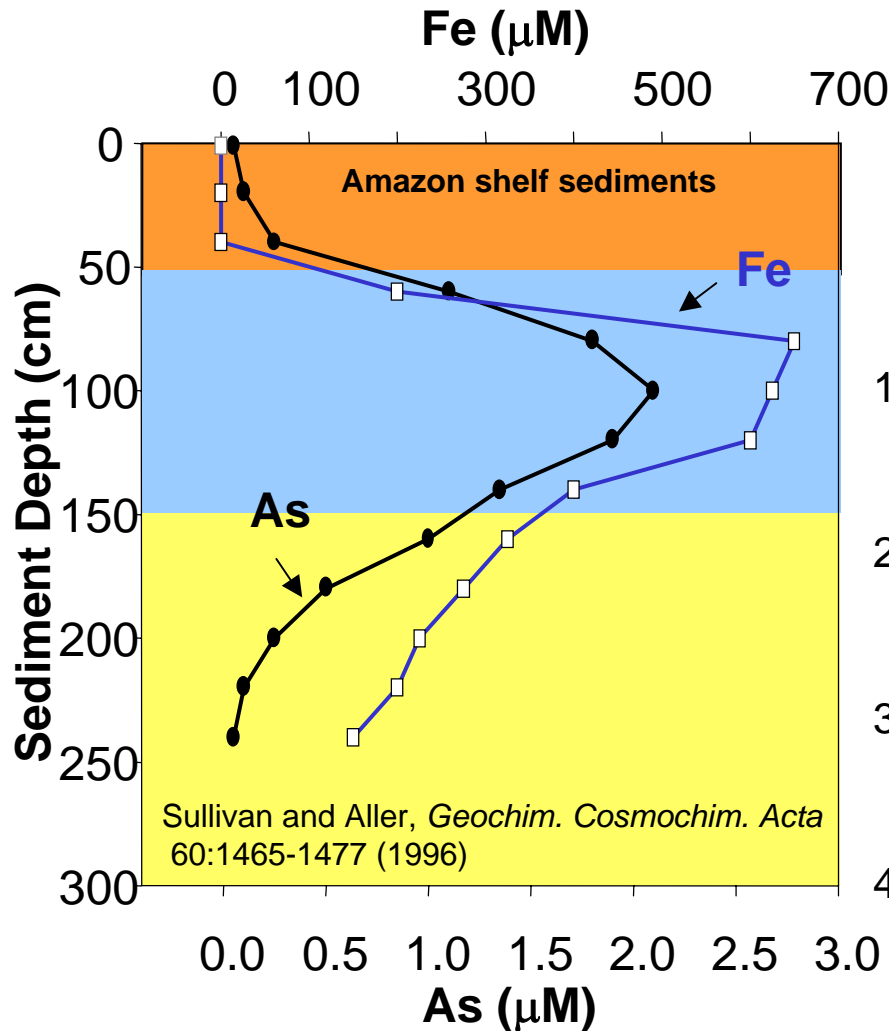
OVER

What Controls Dissolved Arsenic Concentrations in Wells?



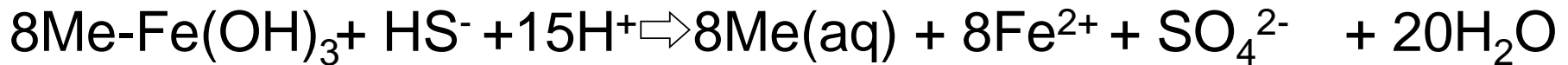
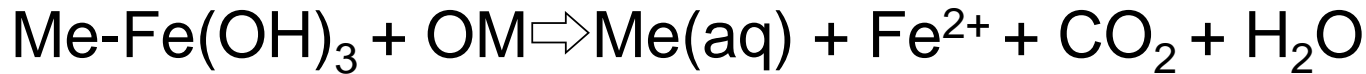
- Arsenic is normally strongly retained by iron minerals
- *Microbes change (metabolize) the minerals in the soil and sediment, thereby releasing arsenic into groundwater.*
- *Conditions usually are reducing (usually +100 to -100 mV) where dissolved arsenic is found.*
- *Organic carbon quality and content critical to the development of reducing conditions*

Arsenic, Iron, and Sulfur Cycling



Trace Metal Retention and Release

Reduction:



However...



Arsenic Sequestration and Mobilization in Model Systems

- Oxidic systems: Fe(III) oxides and sulfate
- Suboxic Systems: Fe(III) oxides \rightleftharpoons Fe(II)_{aq}, sulfate
- Anoxic Systems: Sulfate \rightleftharpoons sulfide, possibly Fe(III) oxides \rightleftharpoons Fe(II)_{aq}
- Field-Based Studies of As Cycling



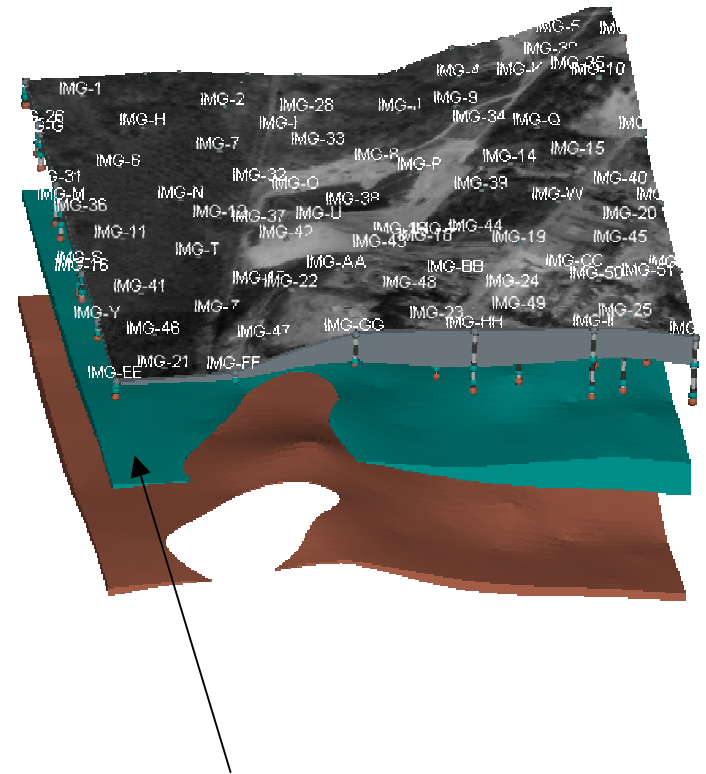
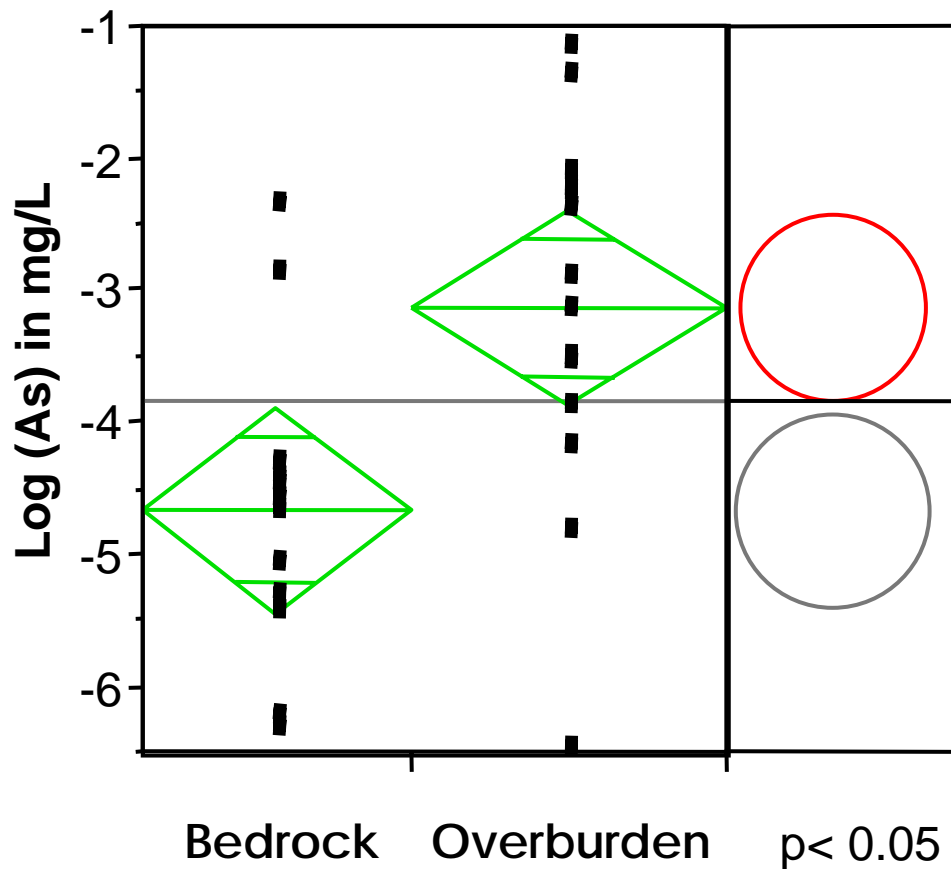
Arsenic-Iron-Sulfur Cycling in 3 Field Sites

- Coakley Superfund Site (NH)
- Coeur d'Alene Mining District (ID)
- Cambodian Groundwater Systems

**Collaborators (Dartmouth): Carl E. Renshaw,
Jamie L. deLemos, Stefan Stürup, Xiaohong Feng**

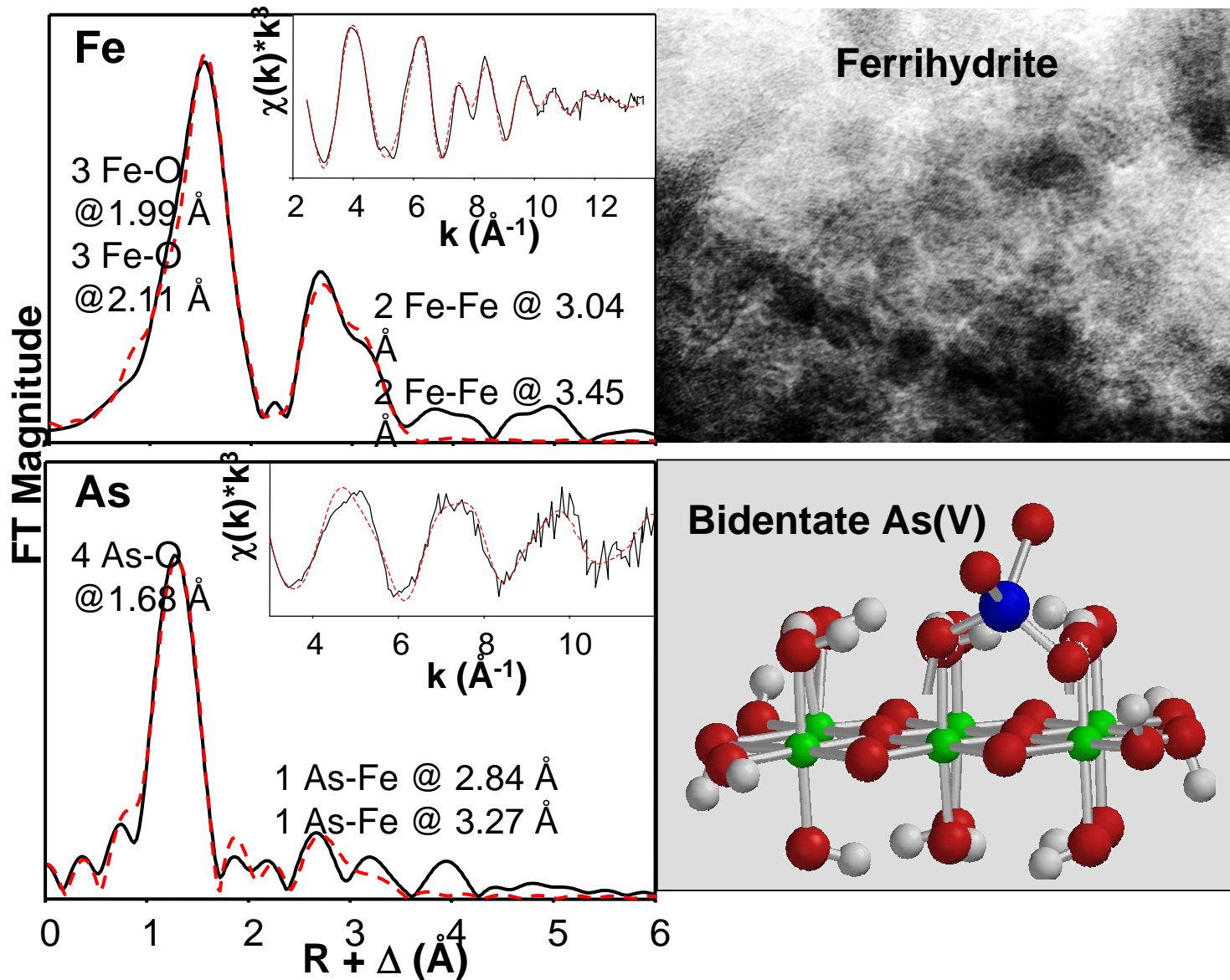
Reference: de Lemos et al. (2005) ES&T

As Source: Overburden-Clay Aquitard

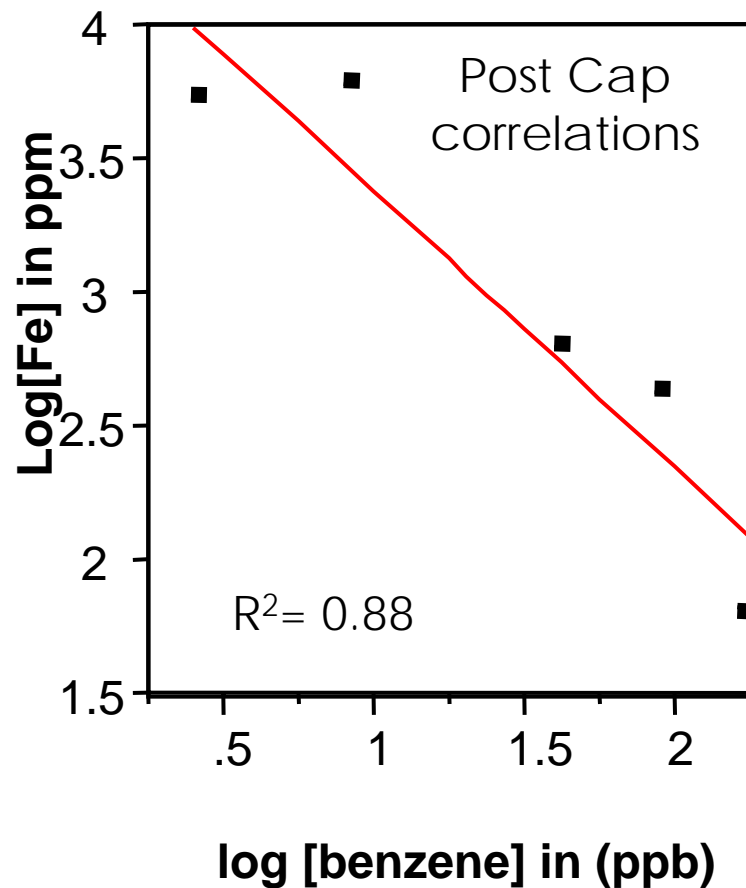
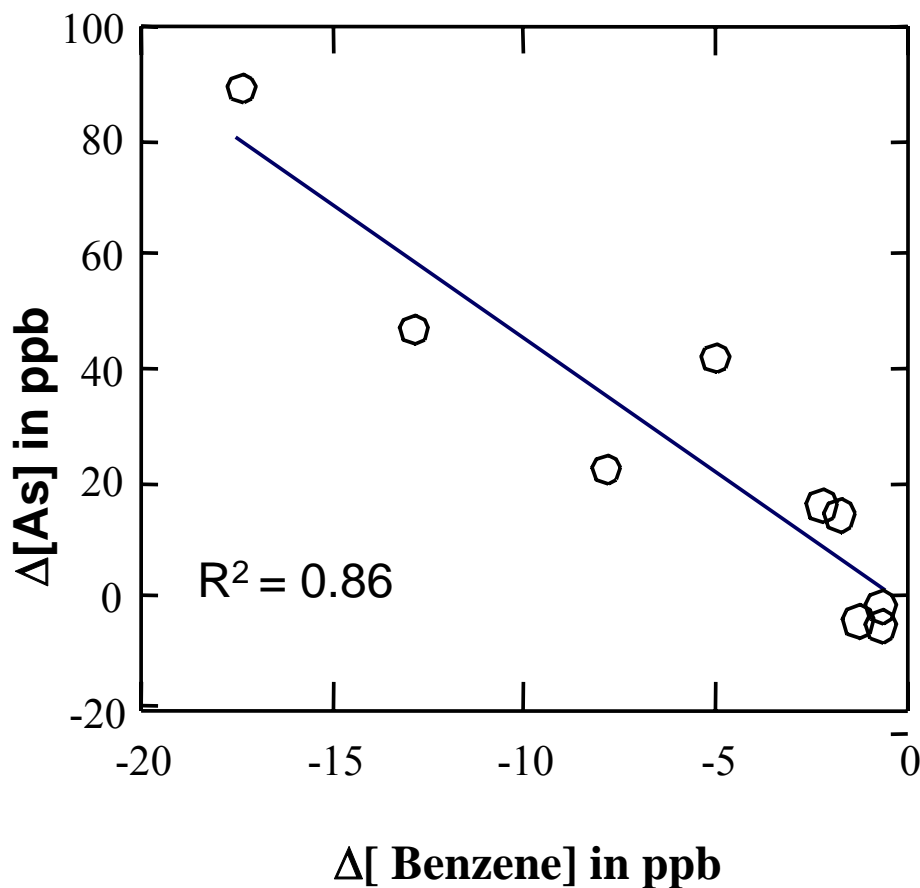
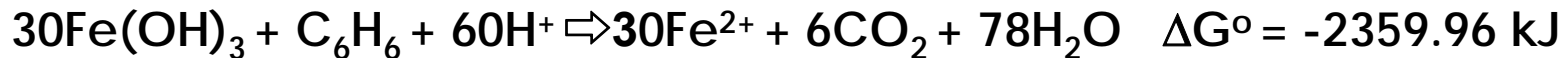


3-10m Thick Clay Layer
 $K = 7 \times 10^{-7} \text{ cm/s}$
[As] ~ 20 ppm

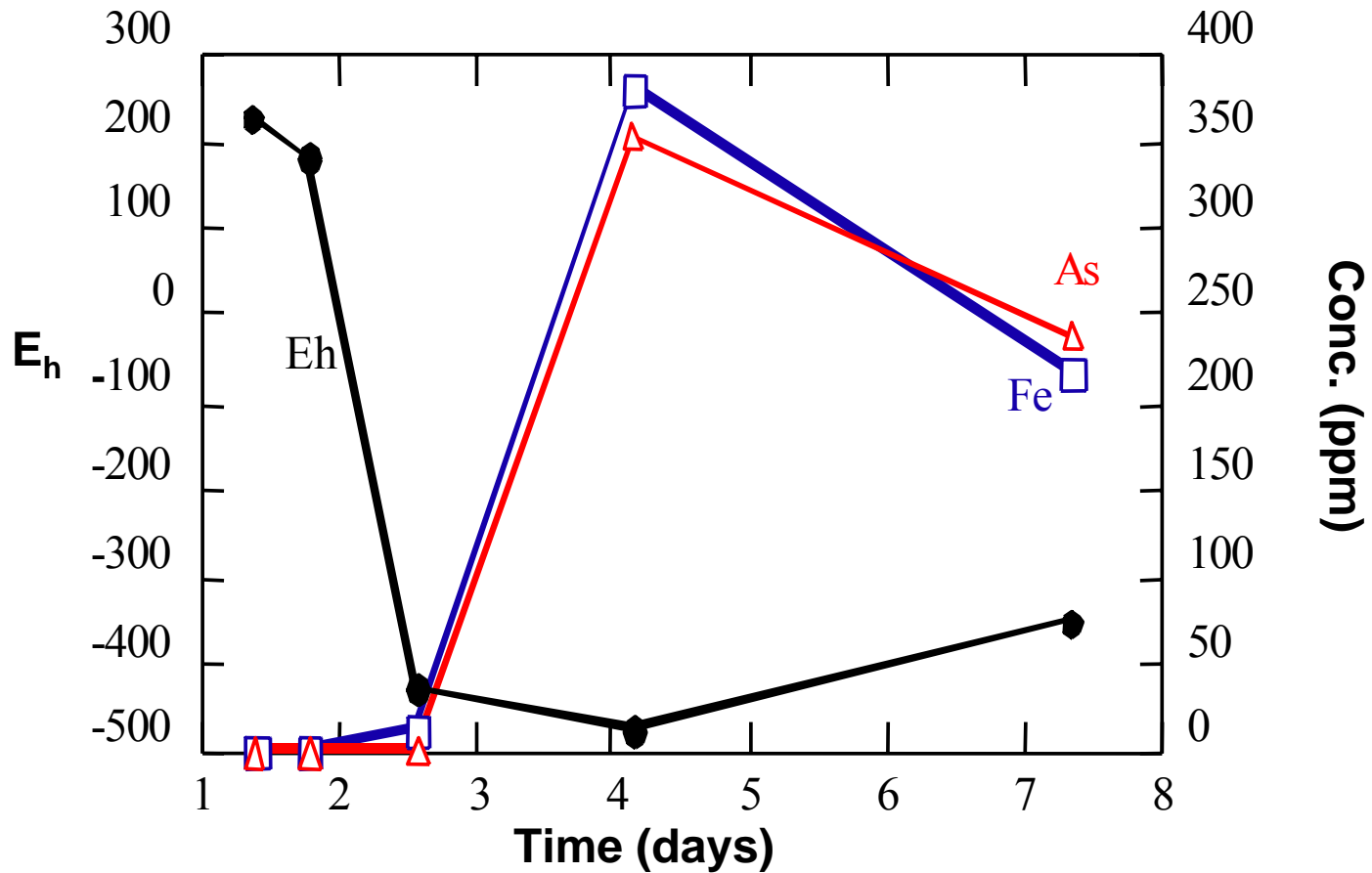
As Source : Characterization



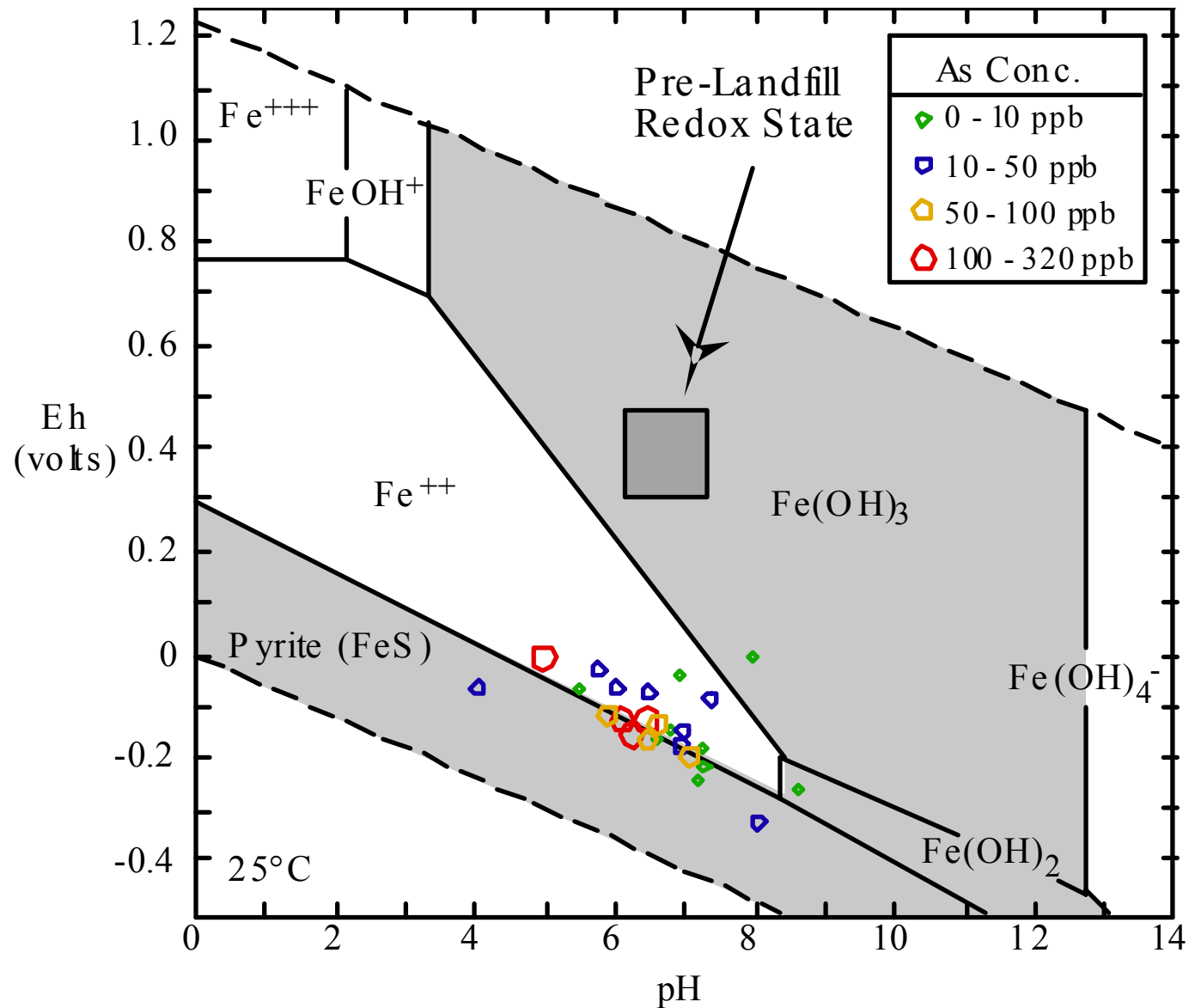
Coakley: Arsenic Mobilization & Natural Attenuation



Coakley: Batch Experiments and Field Data



Coakley: Batch Experiments and Field Data

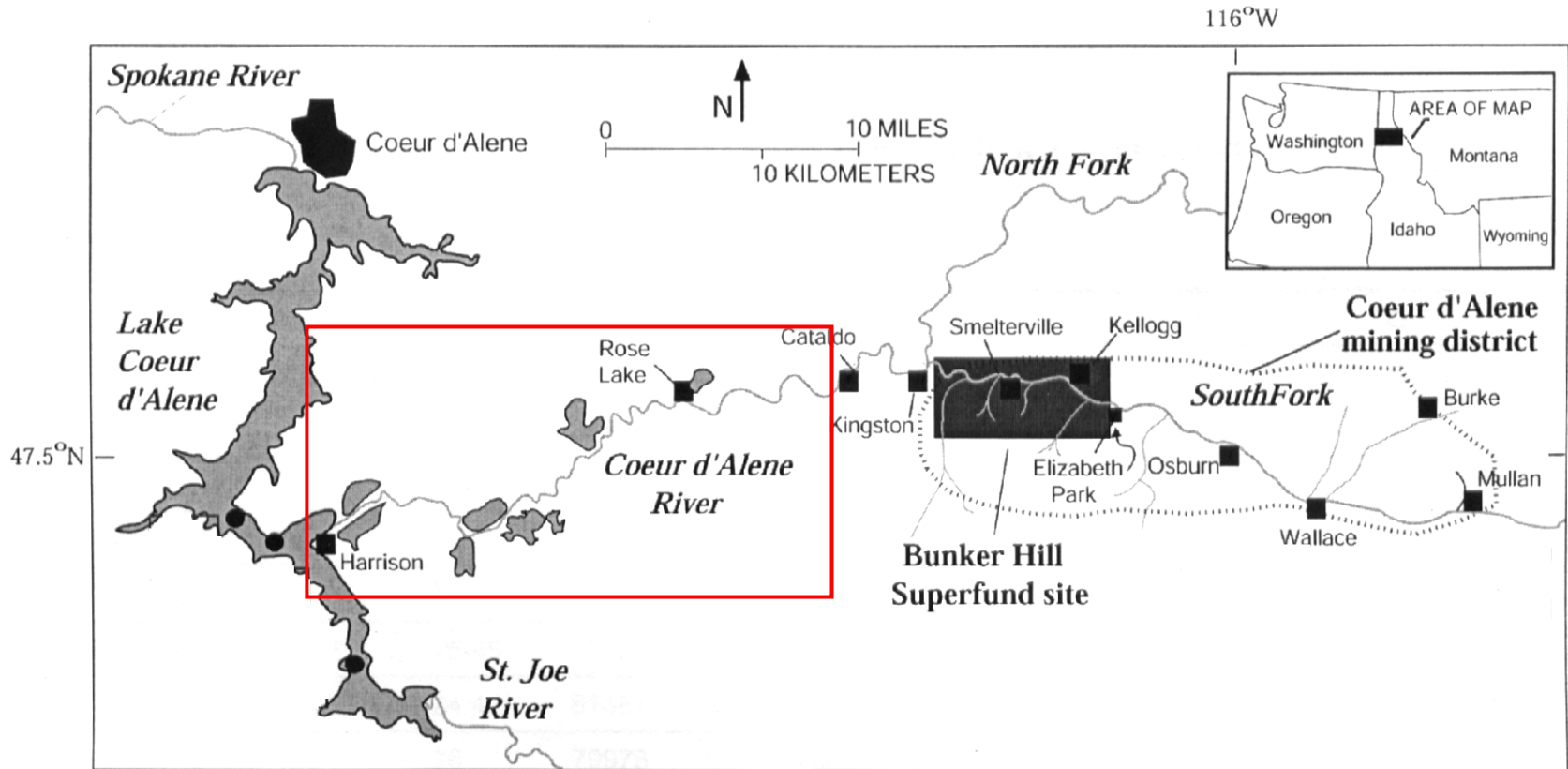


Arsenic-Iron-Sulfur Cycling in 3 Field Sites

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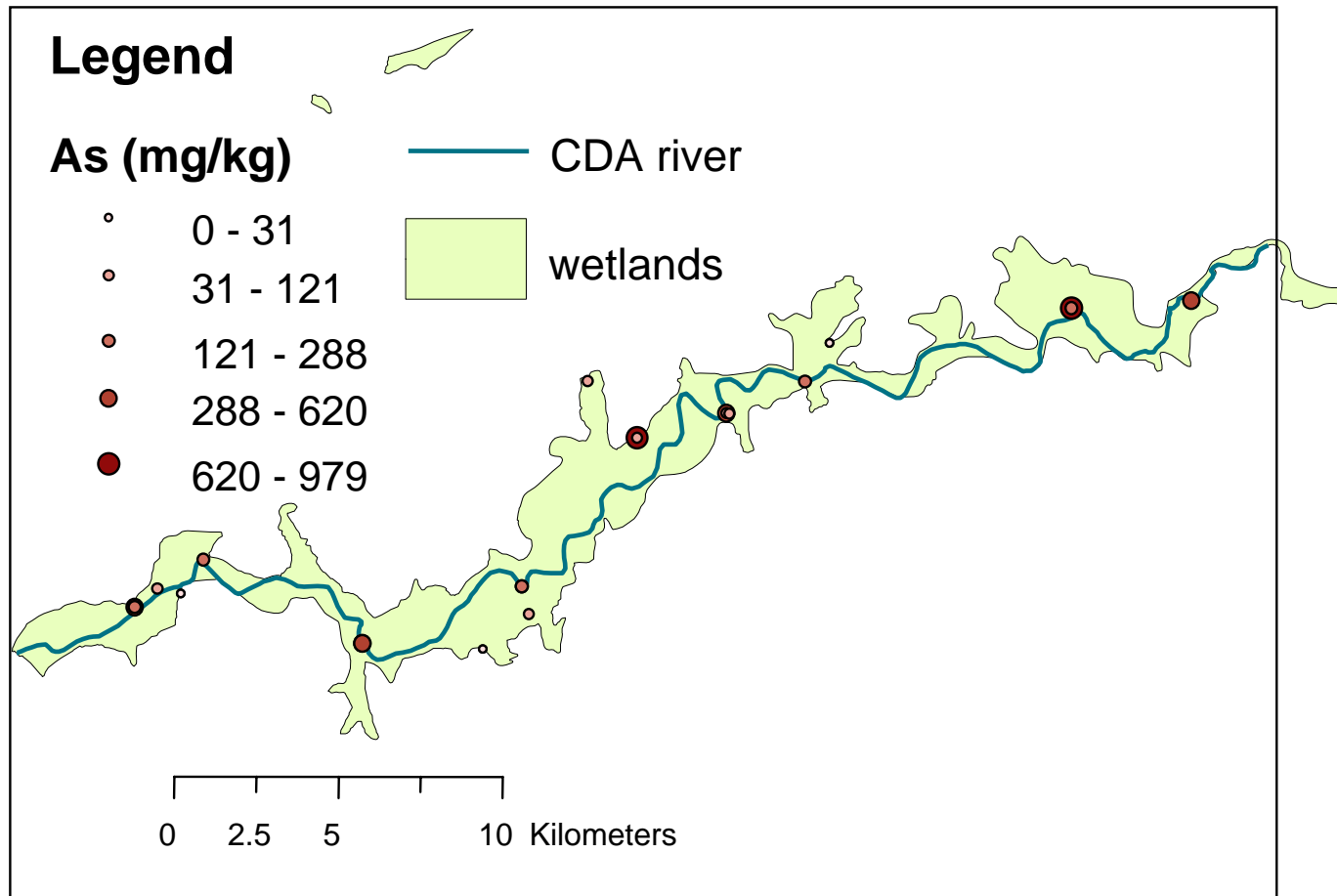
Collaborators: Gretchen Gehrke (Dartmouth),
Gordon Toeve and Matt Morra (Univ. Idaho)
Scott Fendorf and Matt Polizzotto (Stanford)

Coeur d'Alene (CDA) Mining District, ID

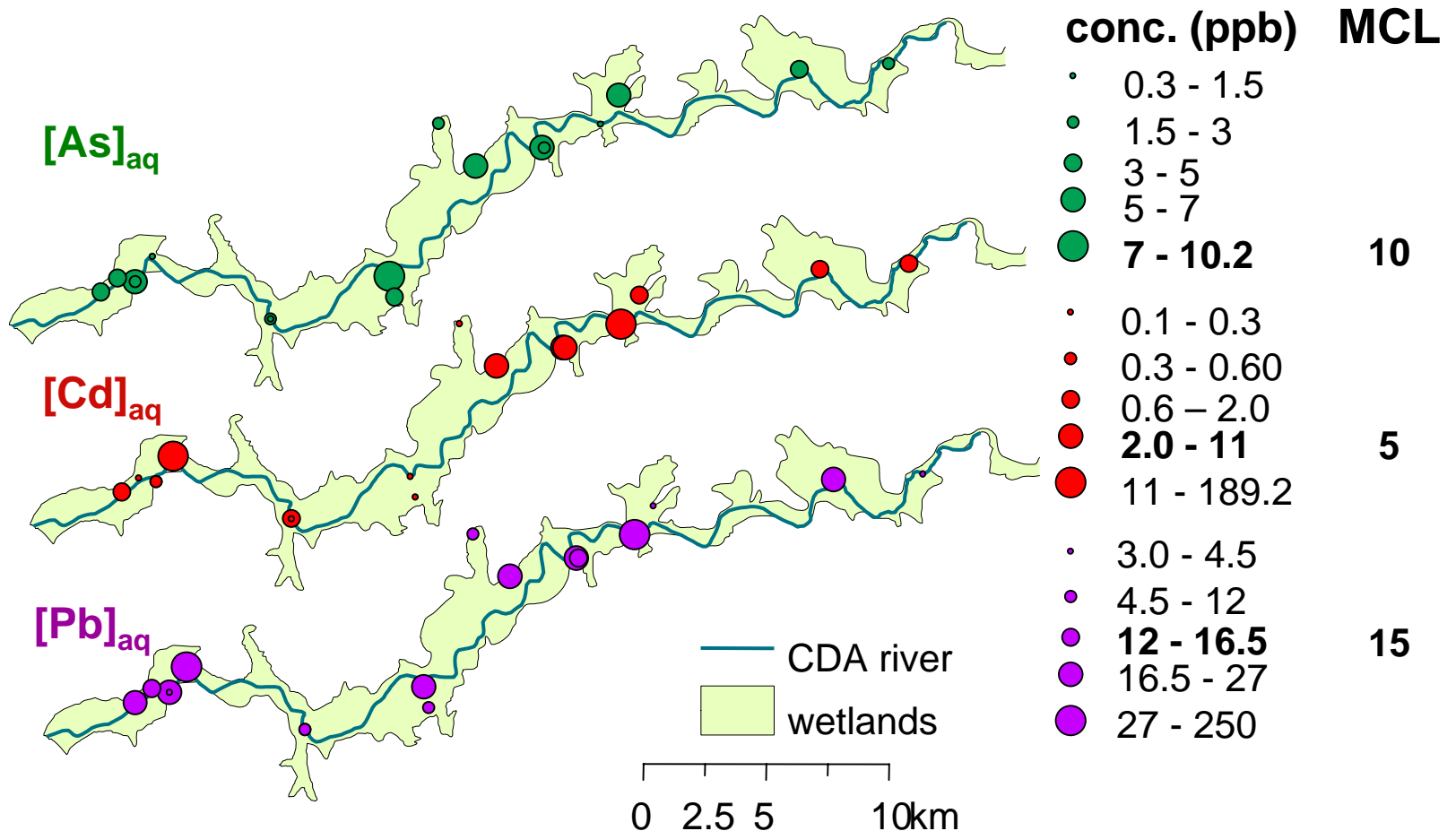


Downstream of the Lateral Lakes

CDA: As Distribution (mg/kg)

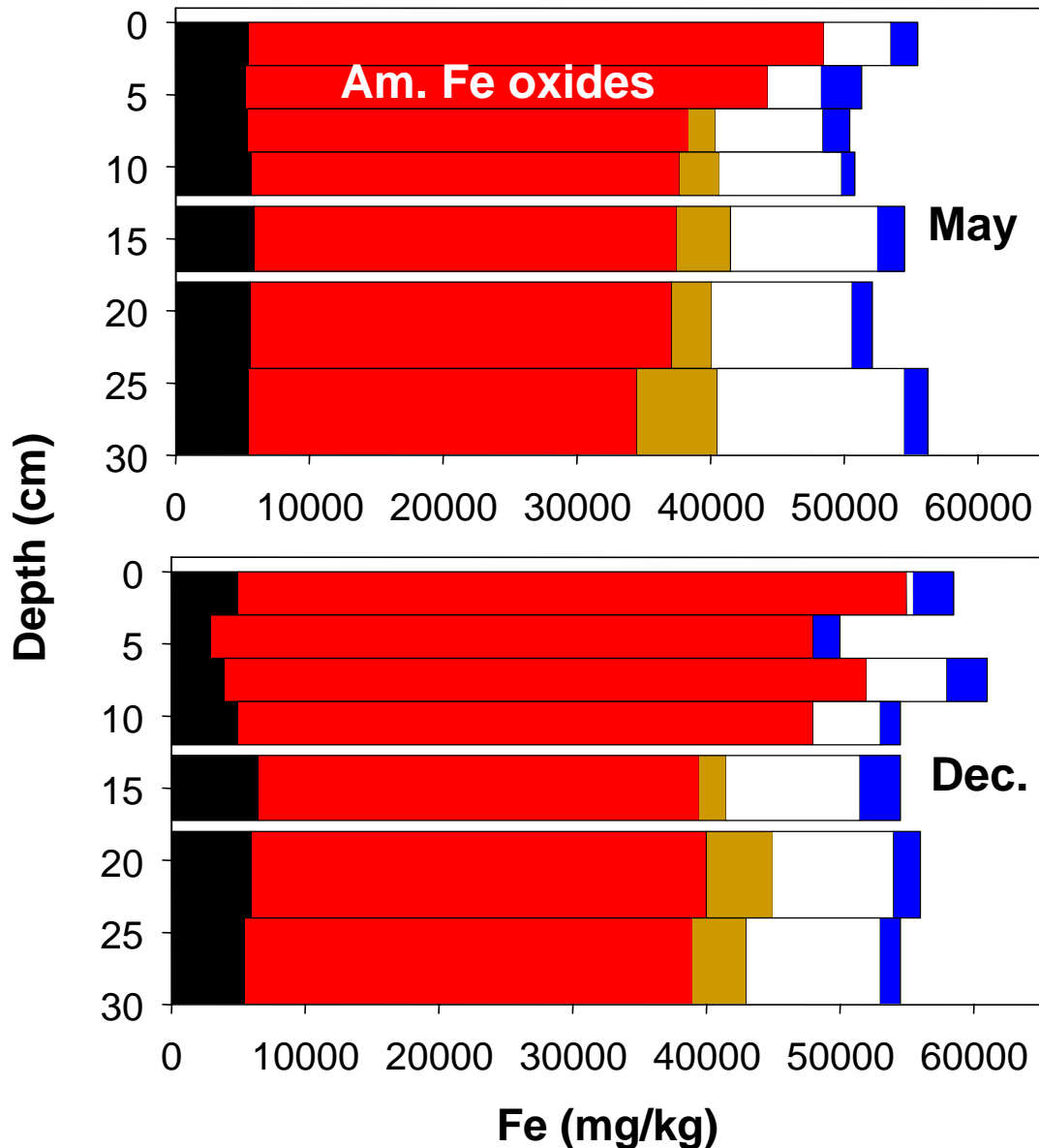


CDA: Dissolved Contaminants

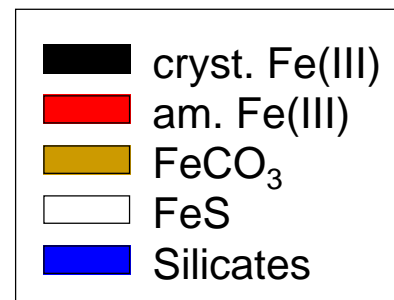


Selected contaminants often are correlated spatially, but in no obvious way with distance from source

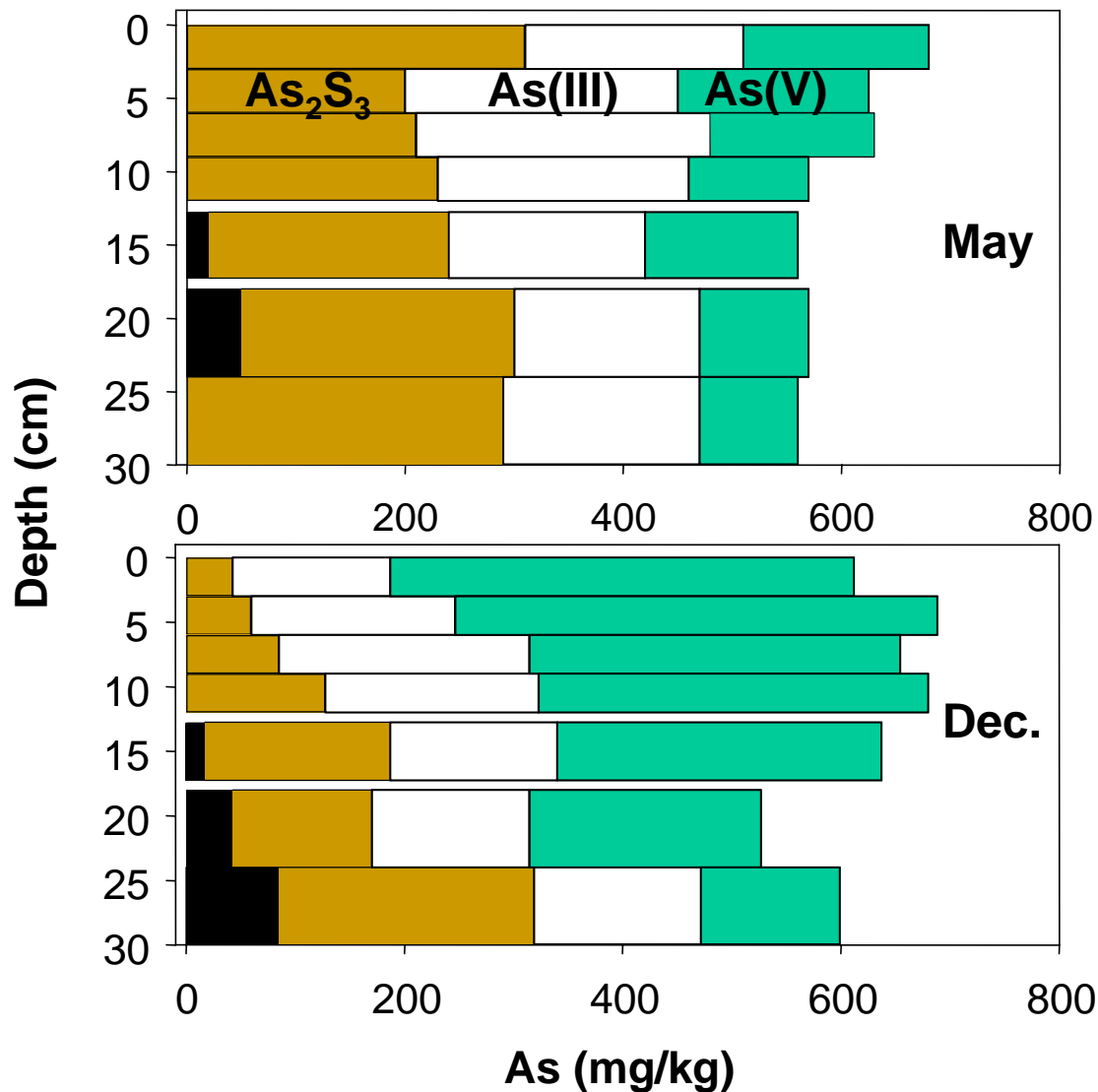
Cataldo: Fe Speciation



- Most Fe is present as amorphous Fe (hydr)oxides
- About 20% maximum fluctuation with season



Cataldo: As Speciation



- Large seasonal variation in the occurrence of reduced arsenic phases in Cataldo Wetland sediments



Experimental Studies of Sulfate Redox Transformations Coupled to As Levels: Coeur d'Alene Mining District

Students:
Andrew N. Quicksall
Samantha Saalfeld
Joshua D. Landis

*Coeur d'Alene River
At Swan Lake, ID*

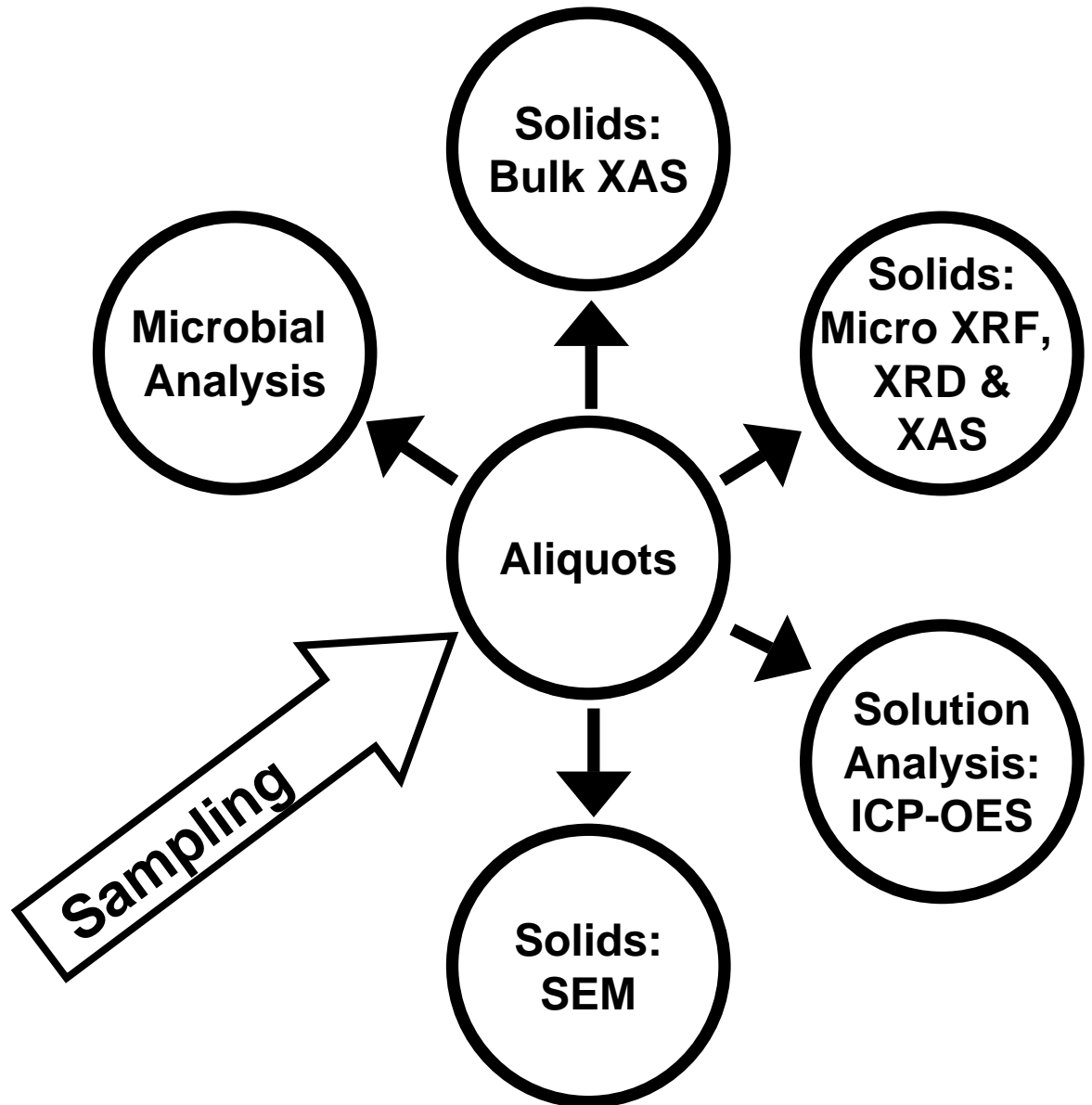
Incubations

Soil Collection

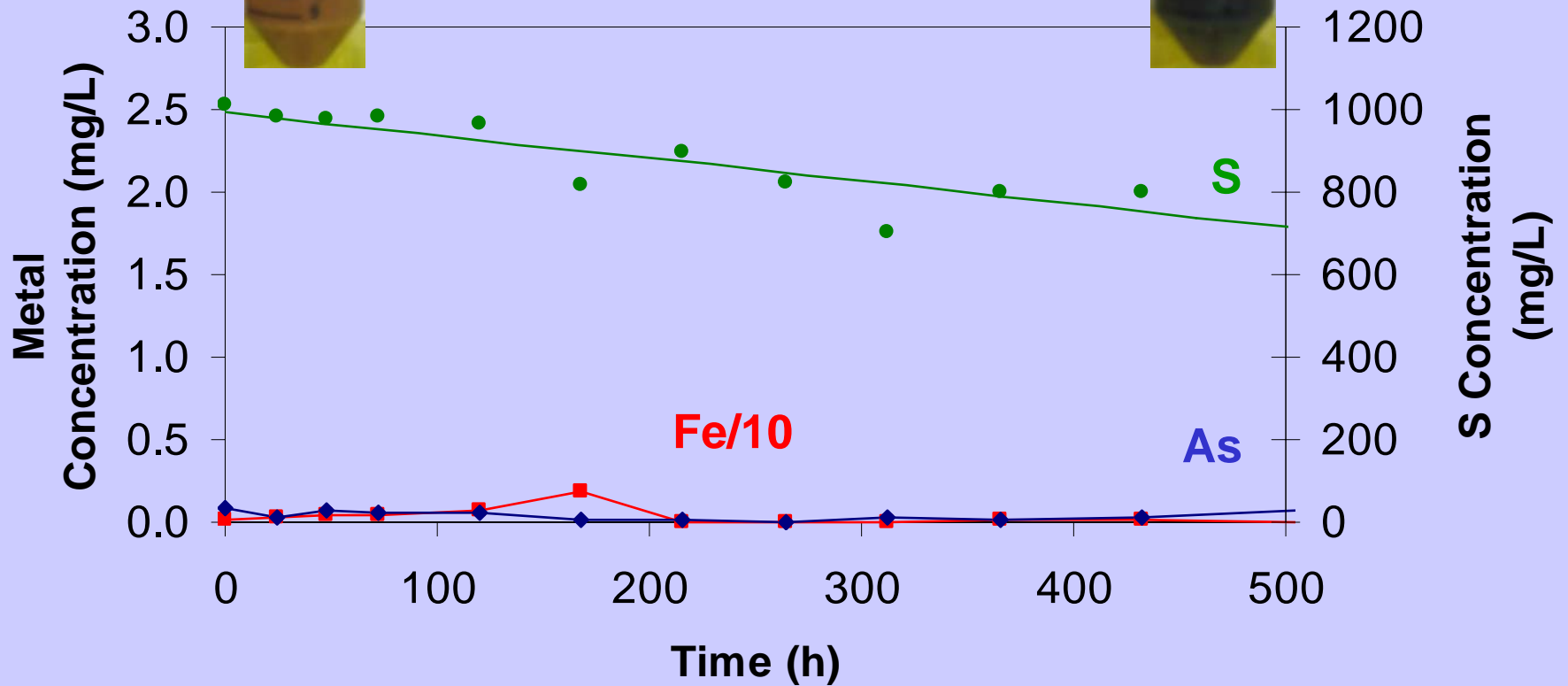


+ Media

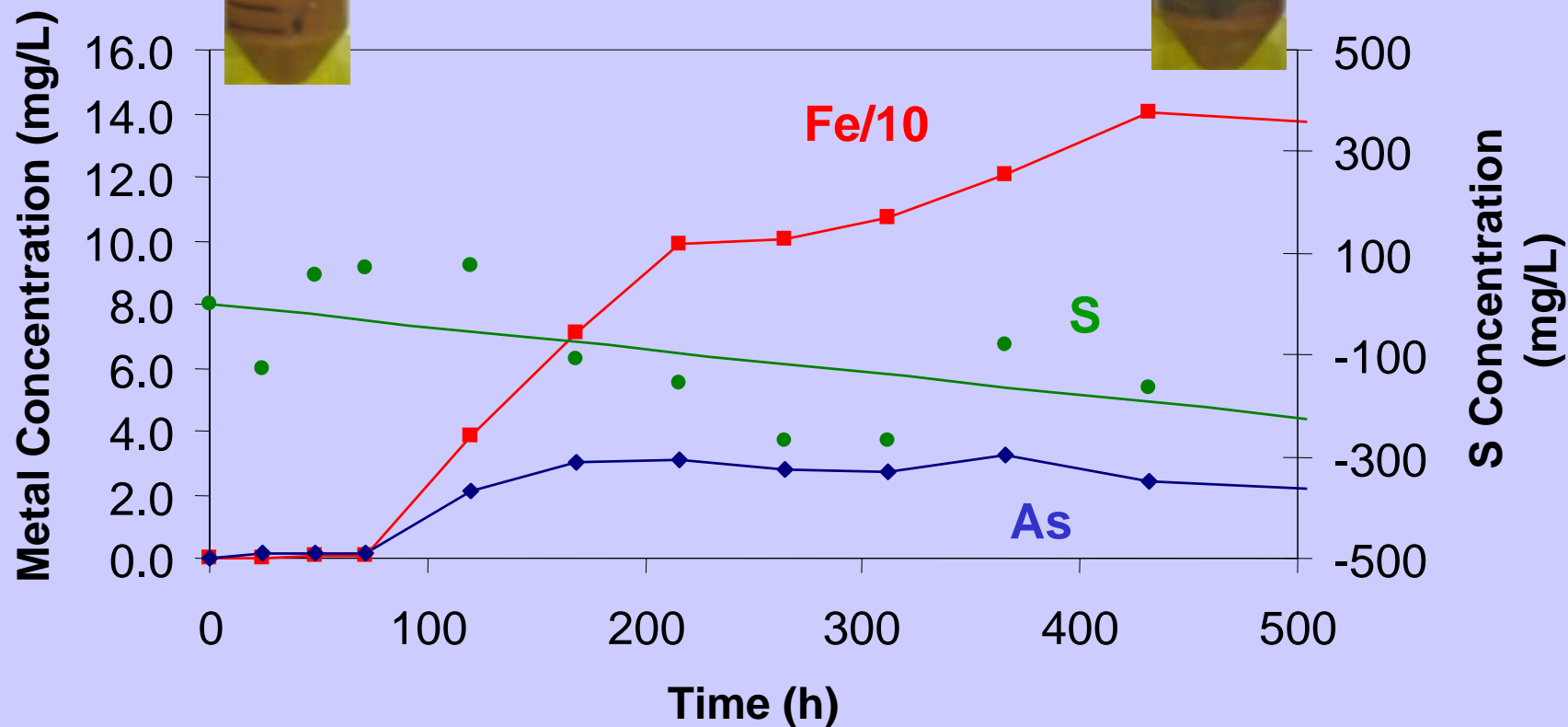
Microcosms



CDA Cataldo Lactate Amended

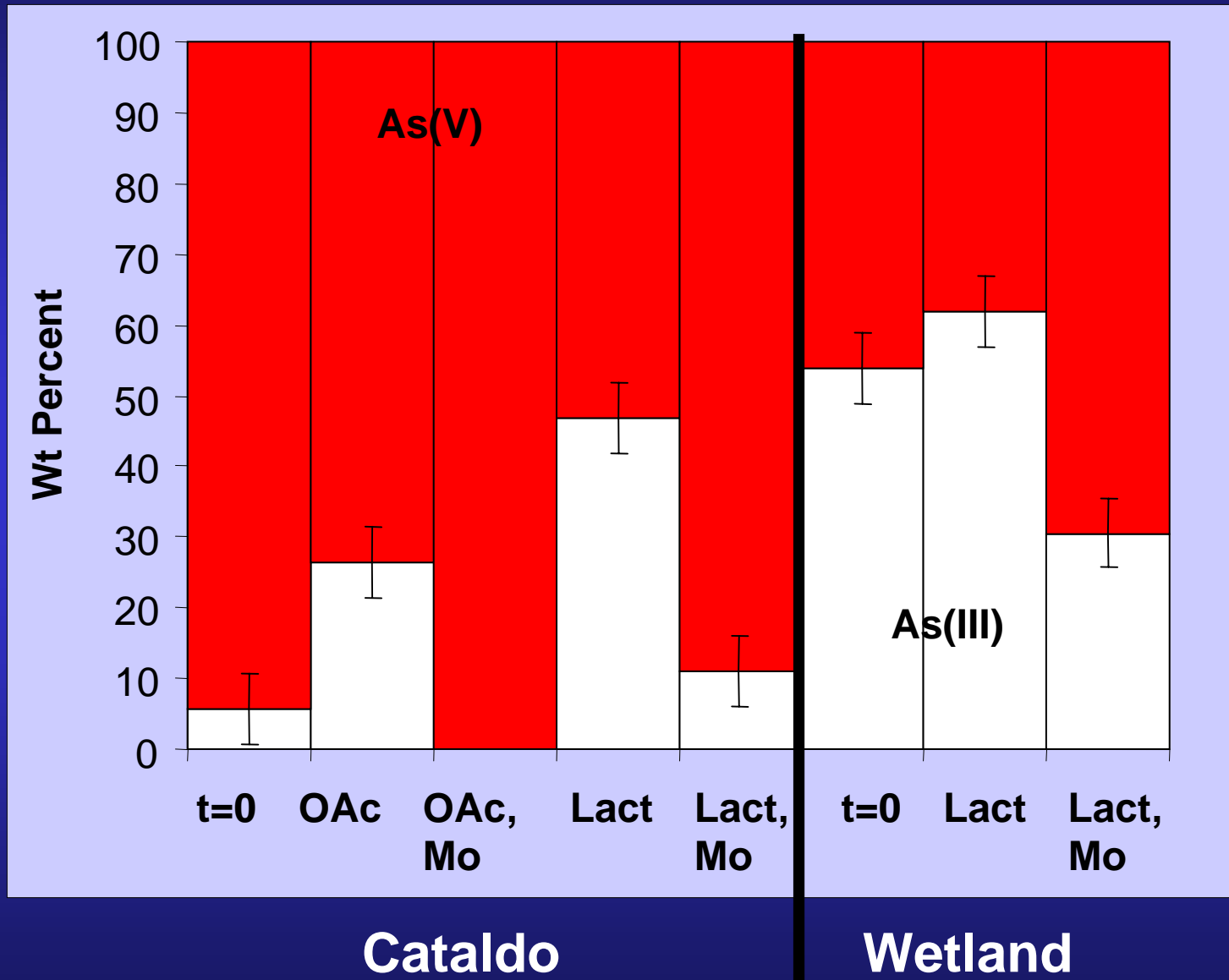


CDA Cataldo Lactate+Molybdate Amended



Plotted as Change from T=0

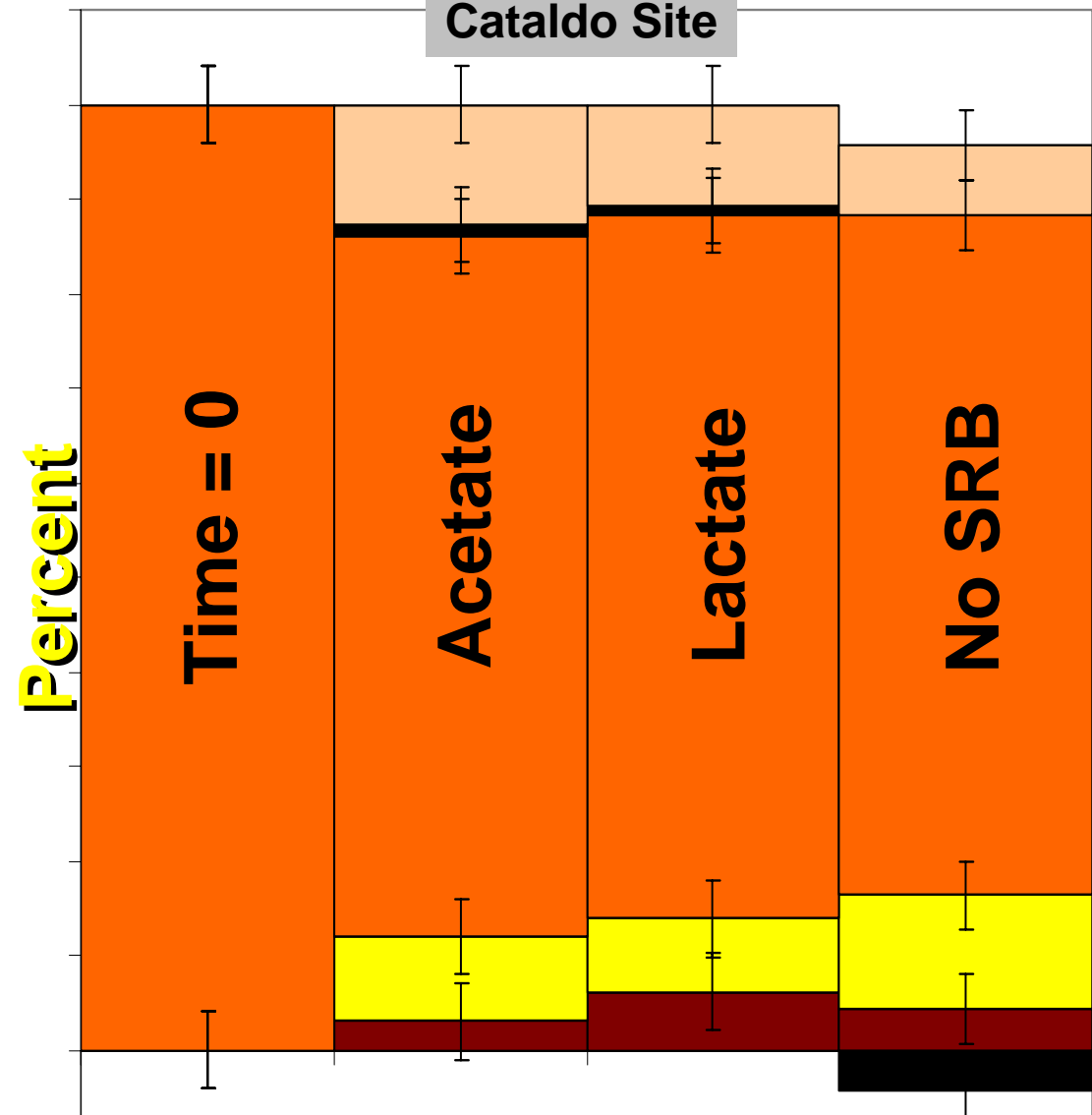
Solids: As Speciation



Determined using As XANES, SSRL 2-3

Original
 Hematite
 FeS
 Siderite
 Goethite

Cataldo Site



Solids: Fe Speciation

Fe(II) mineralization

Siderite present in SRB suppressed with net loss in FeS

Determined using Fe EXAFS, SSRL 2-3

Time=0

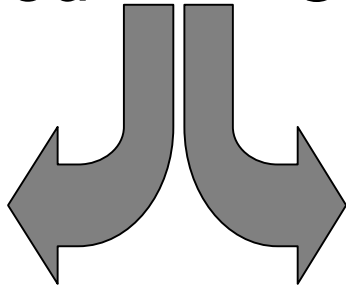


SRB

Full

Suppressed

Community

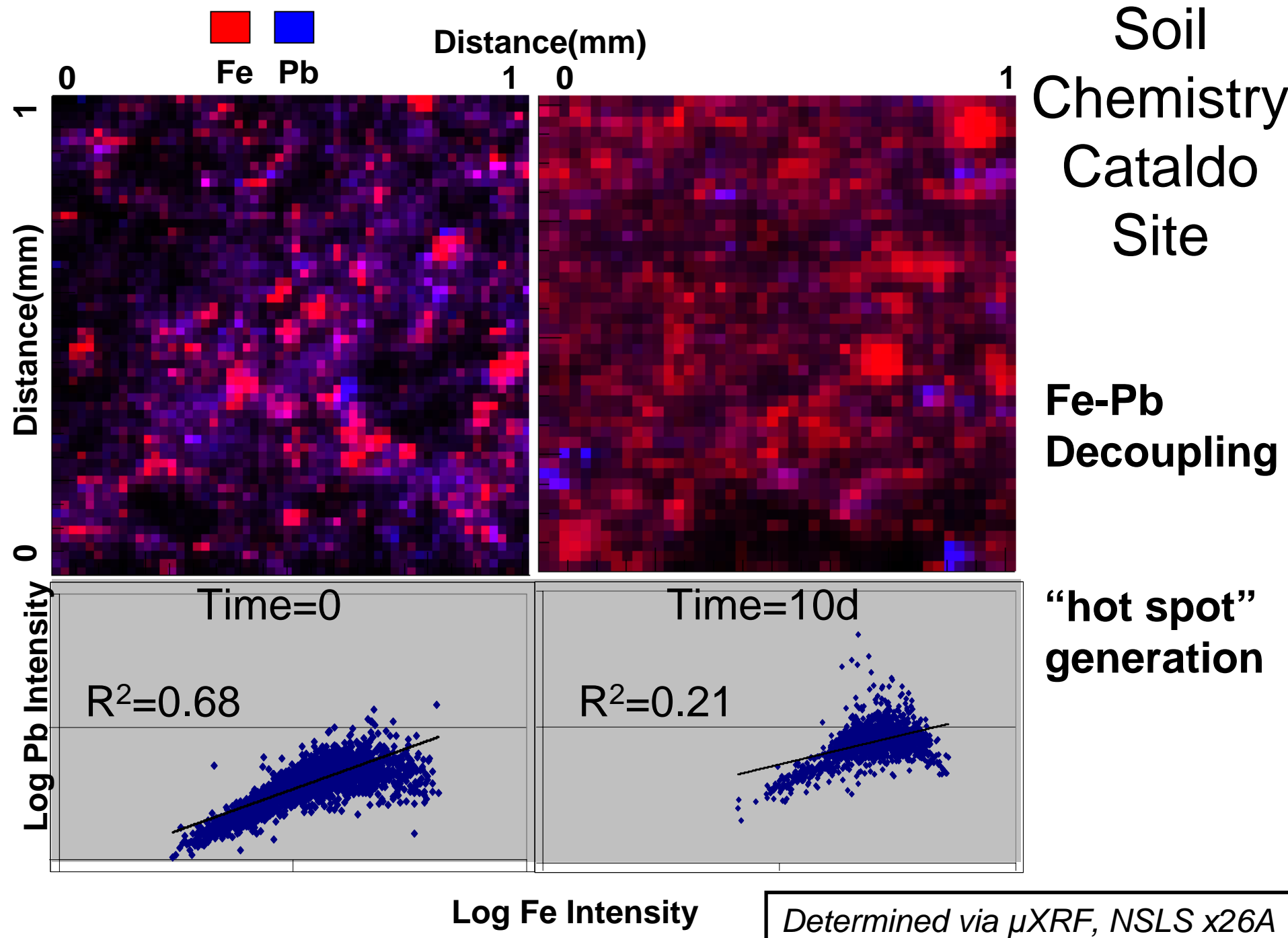


Changes as seen in
representative microcosms

Temporal Change

Paired Fe and As
Release in SRB
Suppressed
Microcosms

Fe and As were
sequestered when
FeRB and SRB were
active

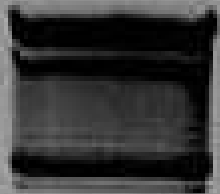


Microbial Ties

- The suppression via molybdate yields strong evidence for SRB involvement in trace element retention
- Can we explain this observation via direct methods to identify specific microbial populations?

Sequencing of cloned 16S rDNA soil extracts

Dominant Microbial Species



Lactate + Molybdate (SRB Suppressed)

Anaeromyxobacter
dehalogenans (100%)

Metal Reducer
Particularly Fe
No S Reducers

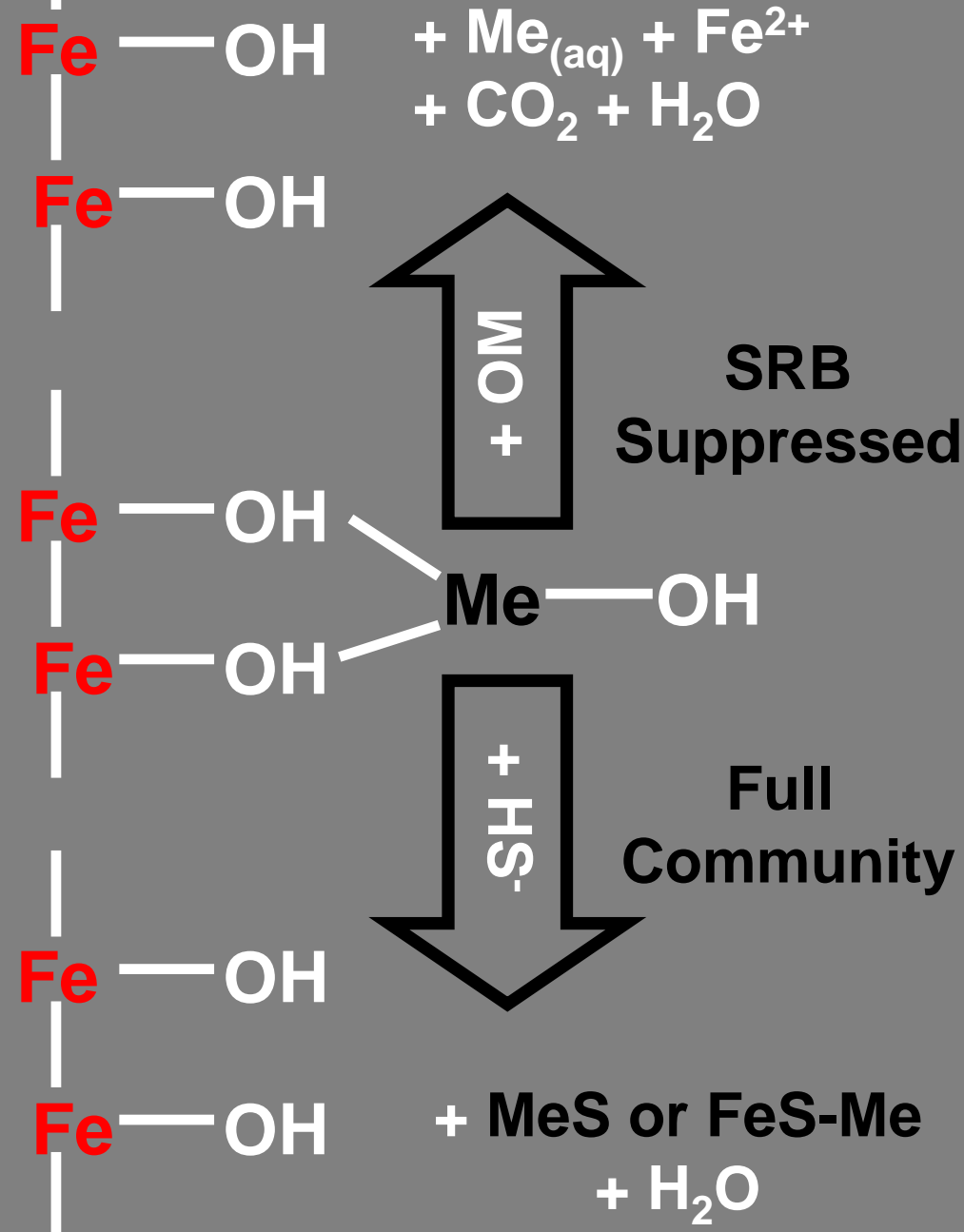
Lactate (Full Community)

Clostridia (89-91%)

Obligate Anaerobes
Many Can Reduce Fe

Desulfitobacterium
hafniense (89%)

Can Reduce SO_4^{2-}



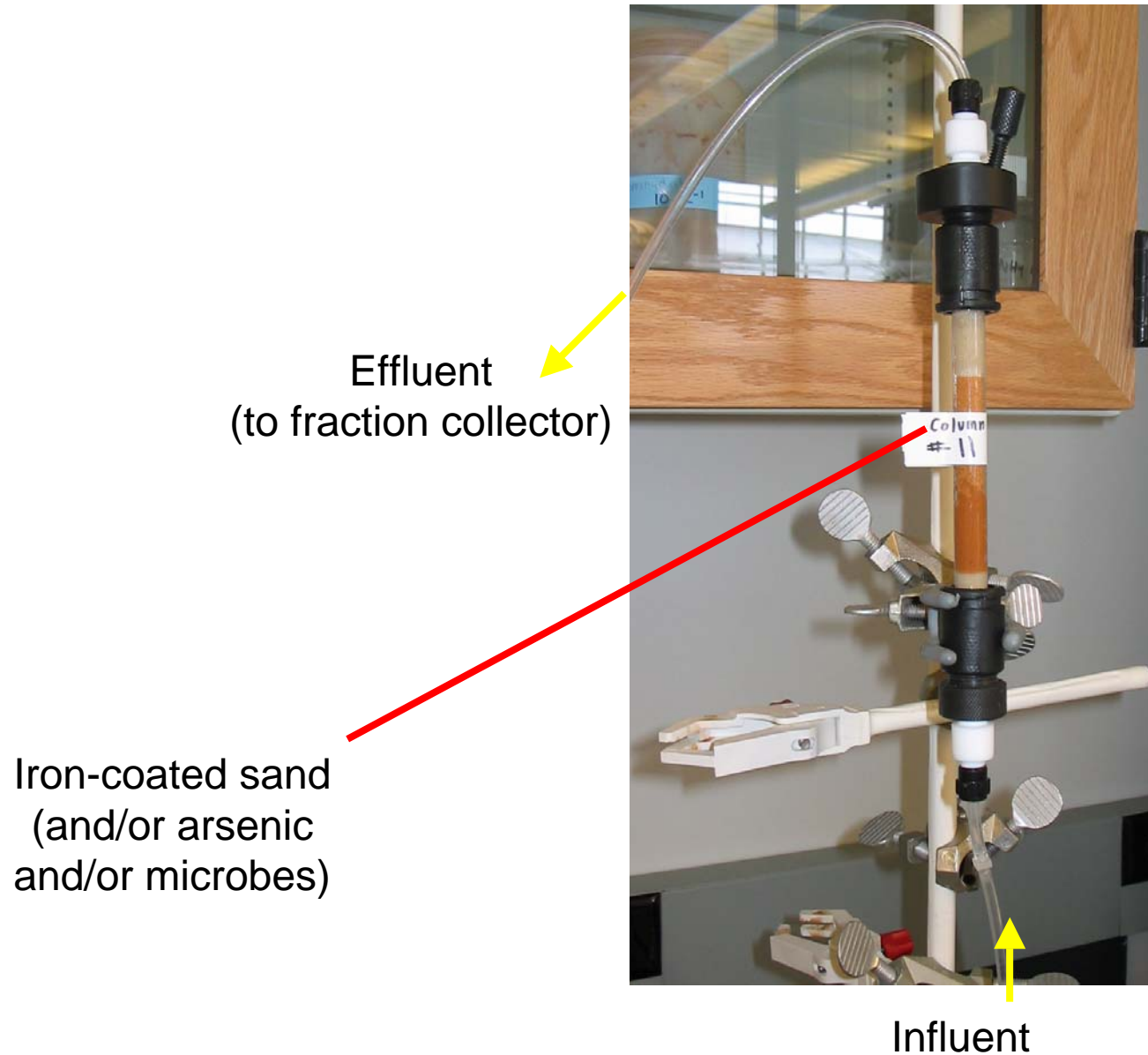
Observations:

- Iron reduction is central to the release of trace metals
- Mineral transformations govern trace metal sequestration

Implication:

- Solution Concentrations are ultimately governed by balanced Fe and S Reduction

Flow-through experiments



Incubations

- Mixed constantly for life of experiment
- Represent stagnant or low-flow end-member of groundwater systems
- Products accumulate, reactants are depleted – system approaches equilibrium

How does sulfate input concentration affect mineralogy in *D. desulfuricans* columns?

5 μ M



0.8 mM



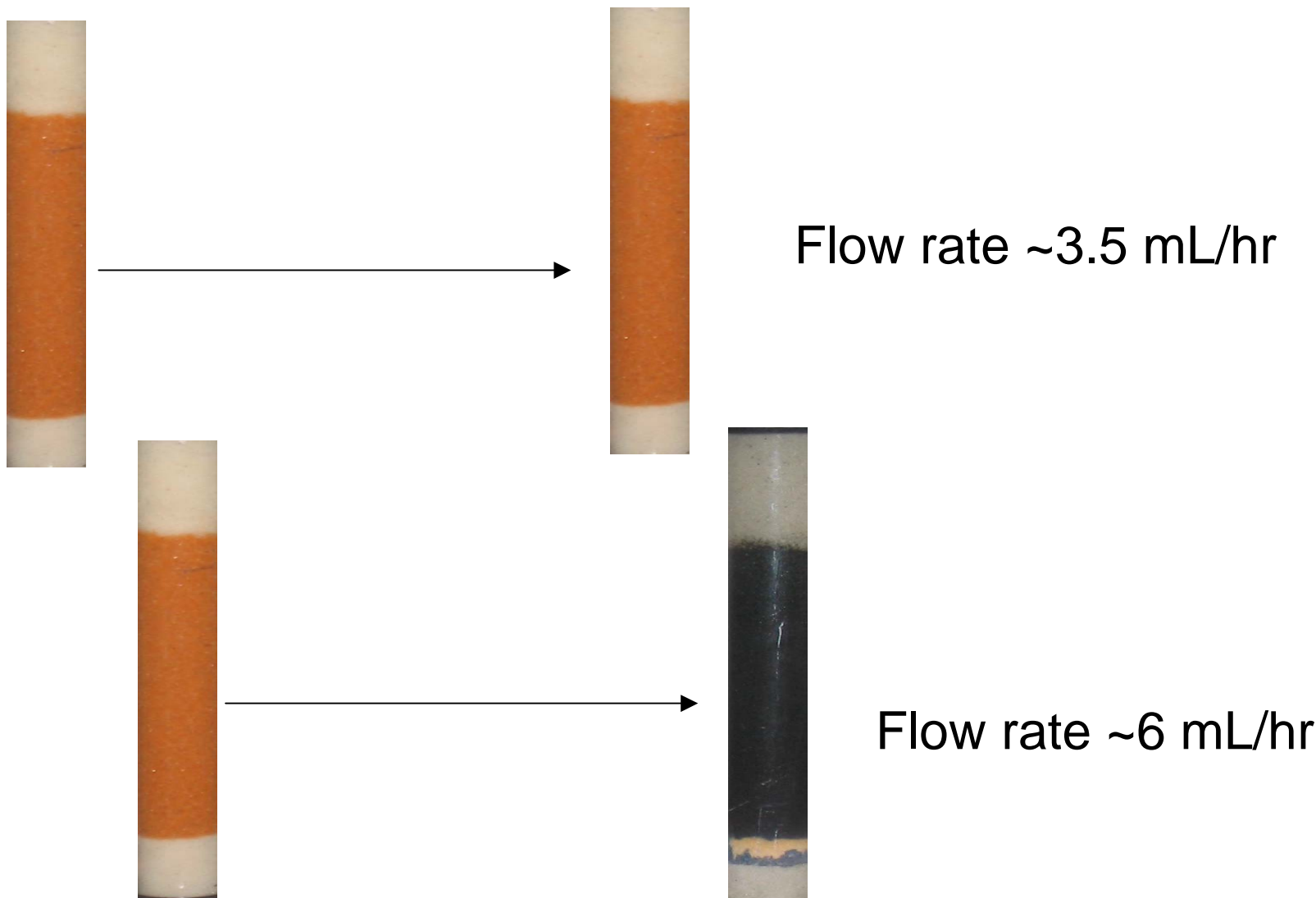
10 mM



20 mM

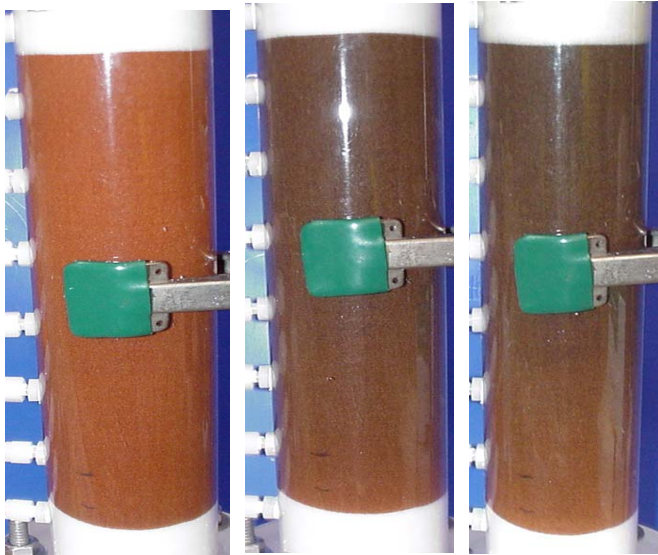


By what mechanisms does flow rate affect mineralogy in iron oxide-sulfide columns?

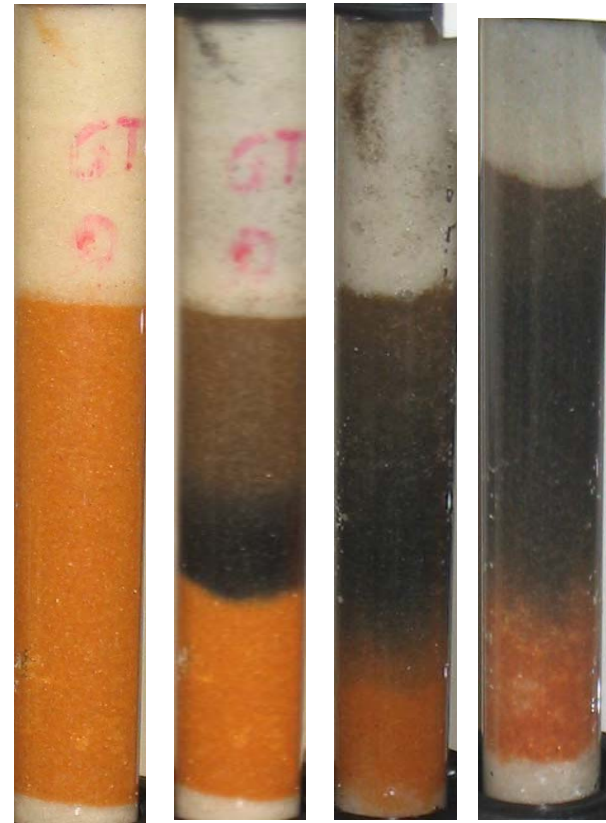


What determines which minerals form in SRB/FeRB systems?

- Magnetite formation



- Magnetite + iron sulfide formation



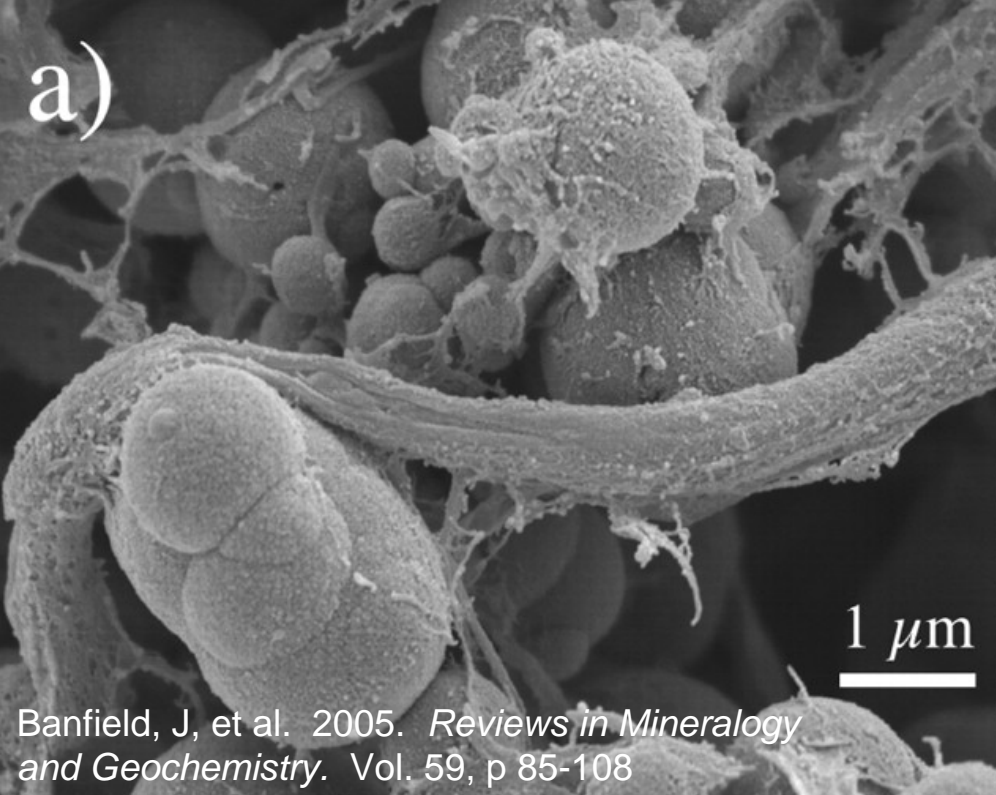
What mineral transformations occur under stagnant conditions in *D. desulfuricans* incubations?

SRBs, 10 mM SO_4

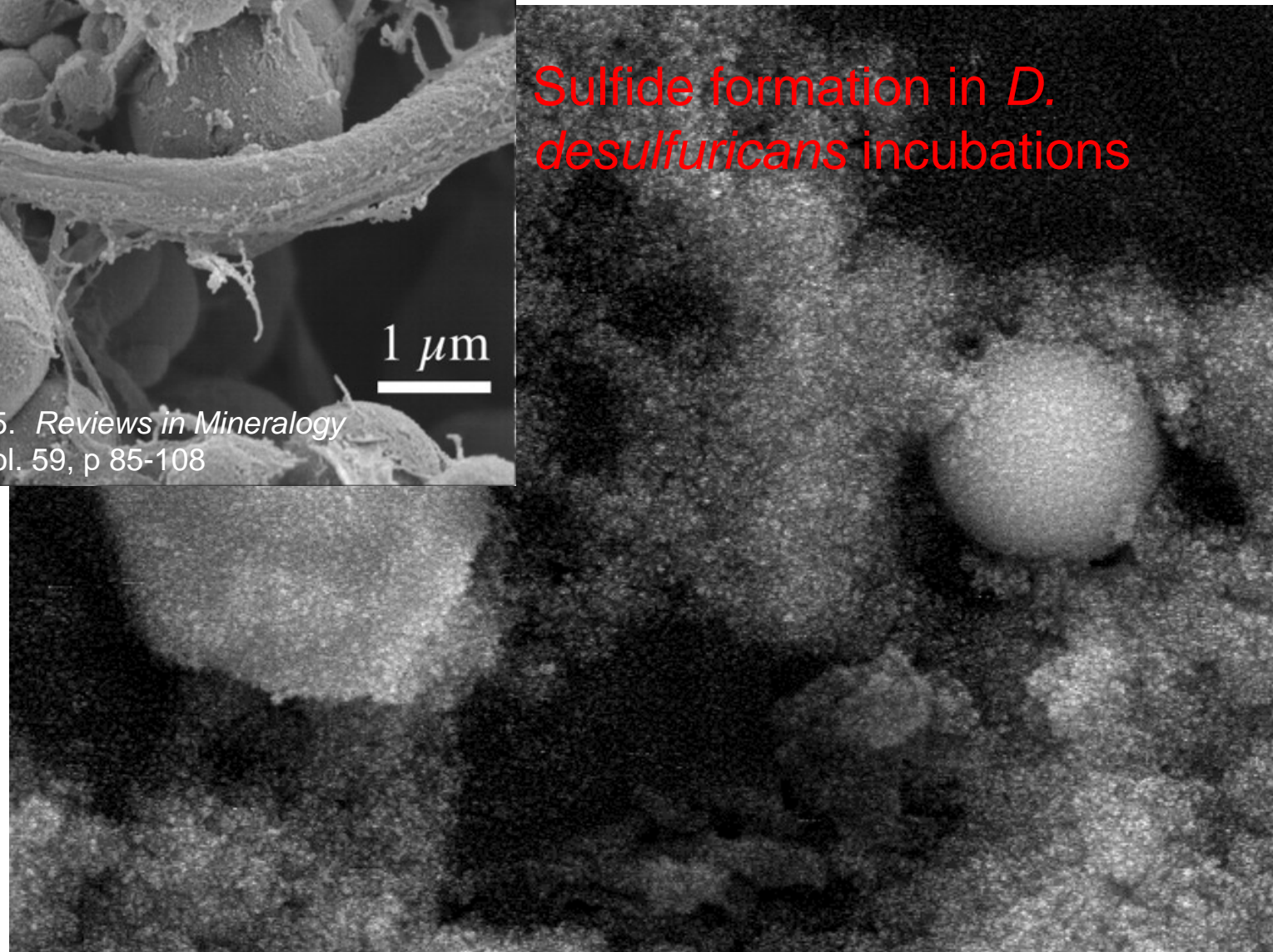
No bugs



**Only modest changes in Fe mineralogy
(not enough carbon to reduce all Fe)**

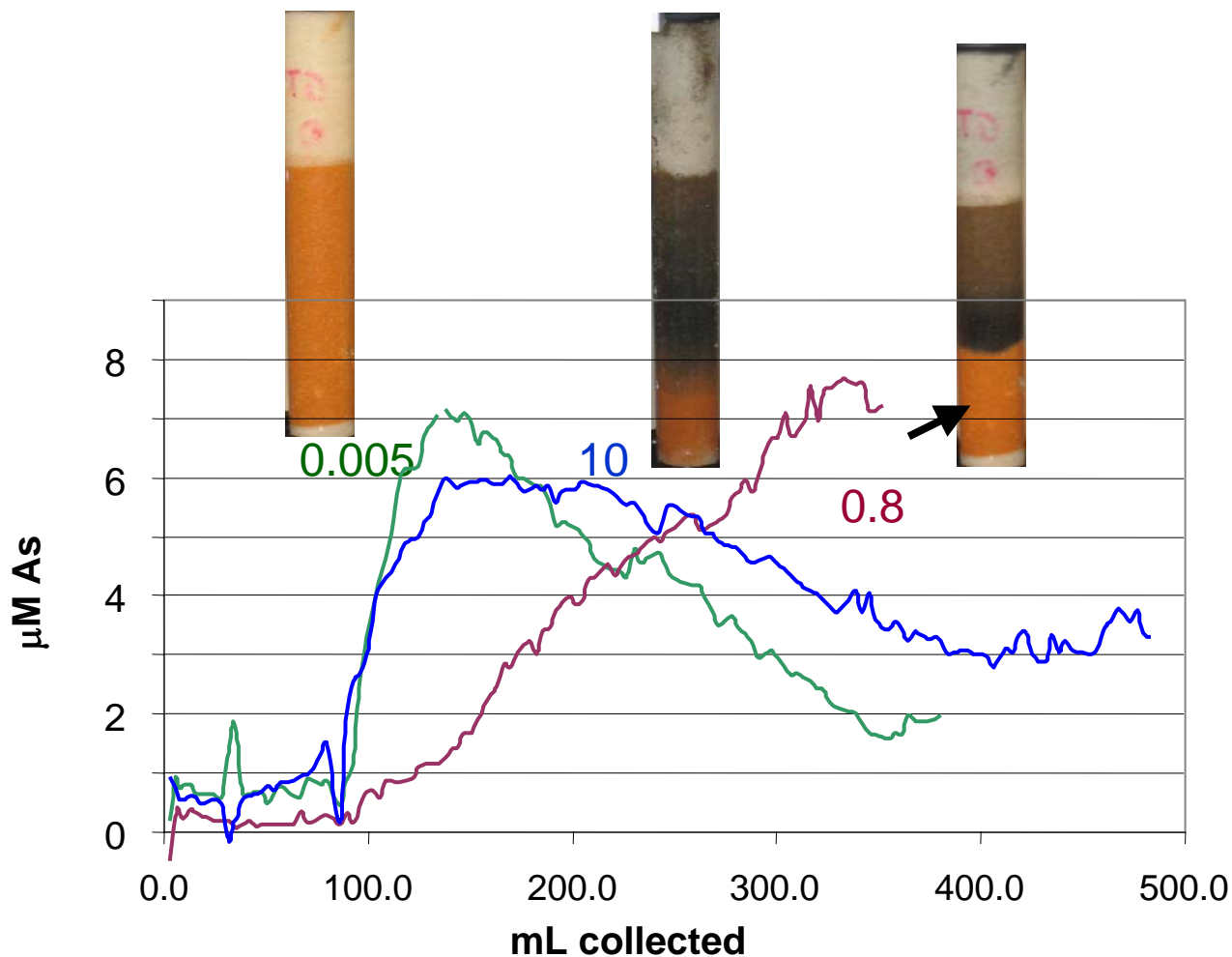


Sulfide formation in *D. desulfuricans* incubations



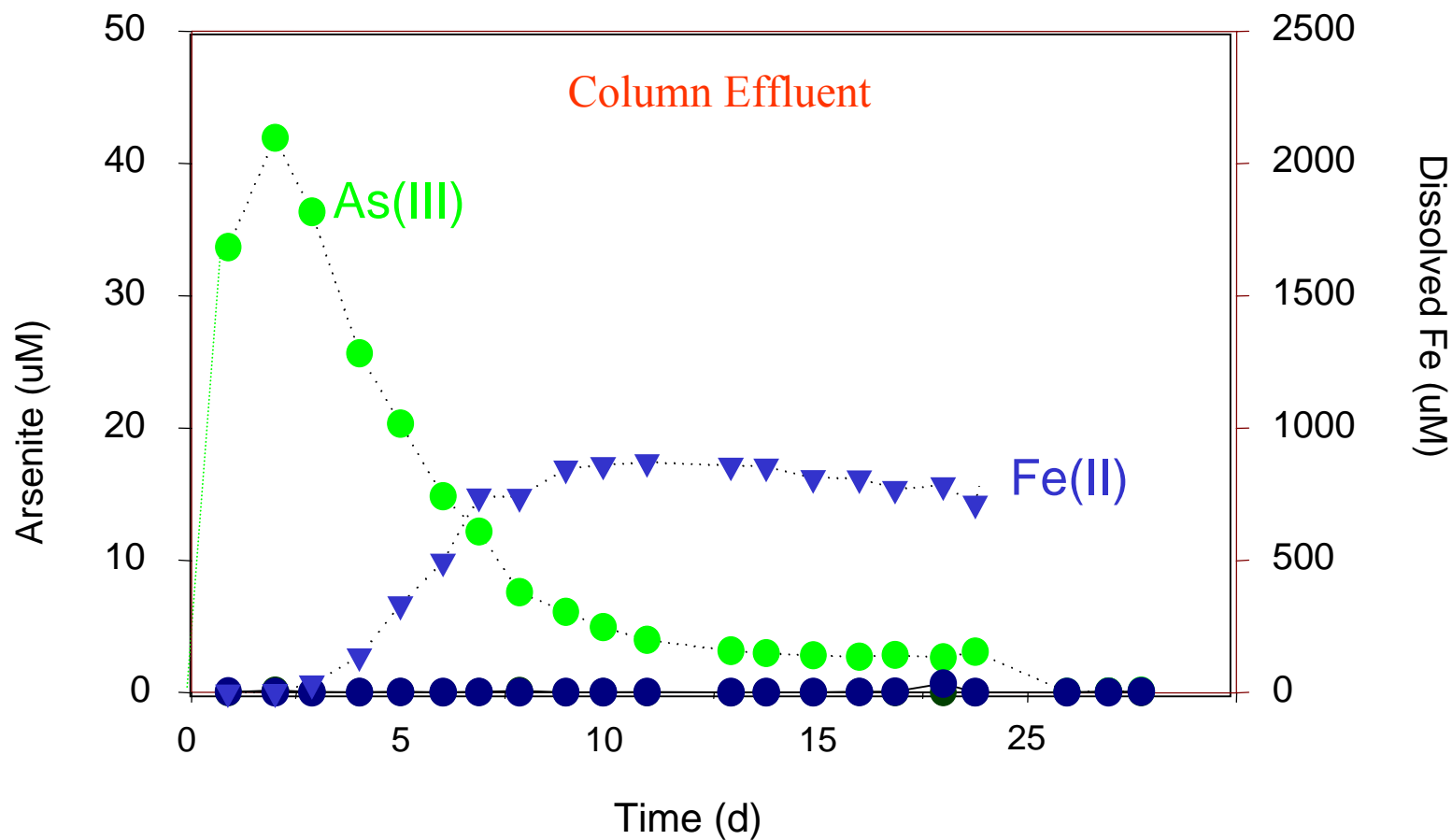
Acc.V Spot Magn Det WD | 2 μm
15.00 kV 2.0 10000x SE 10.5 | Dartmouth E. M. Facility

What mechanisms regulate arsenic release from *D. desulfuricans* columns?



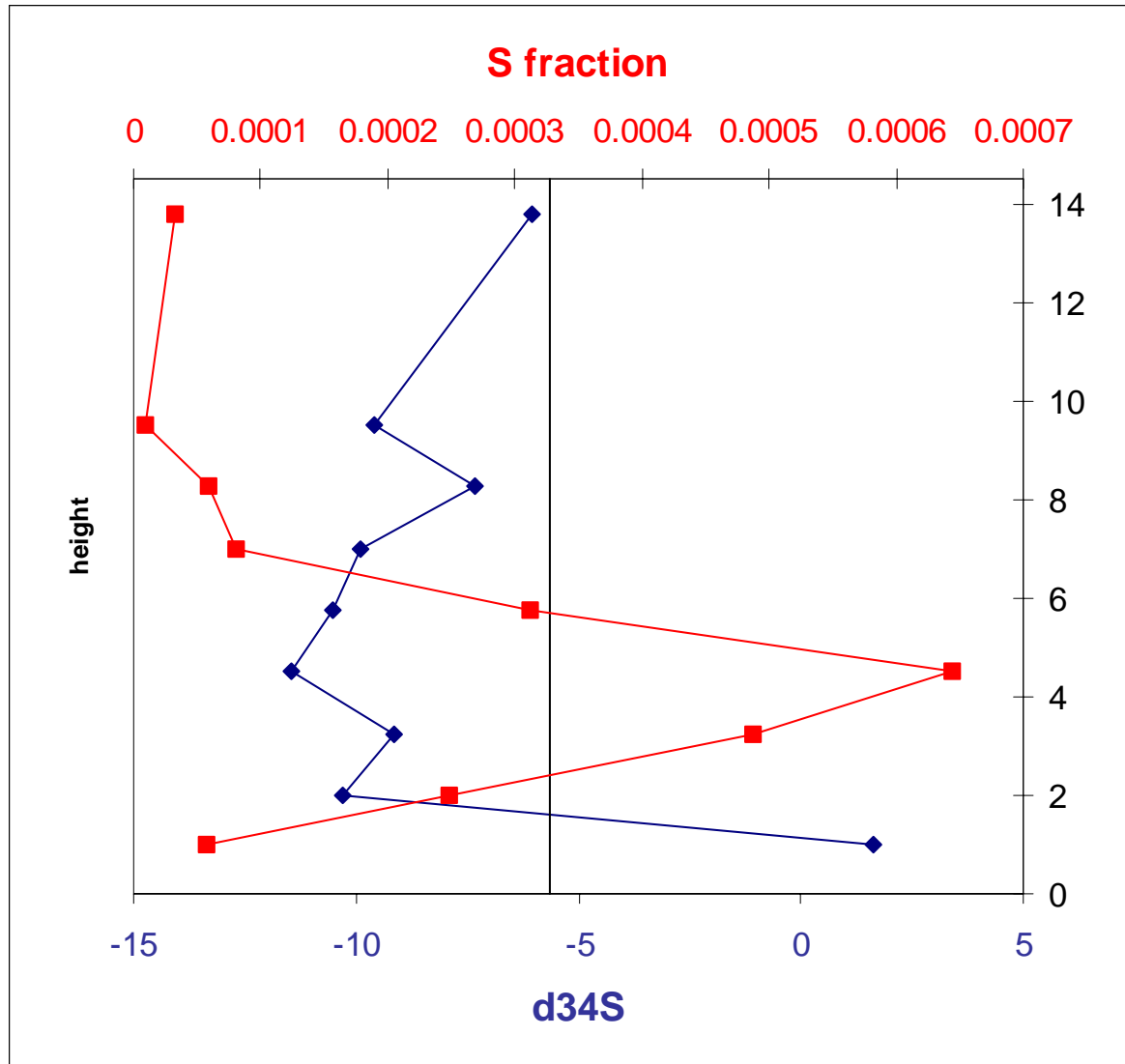
— As (0.005 mM sulfate) — As (0.8 mM sulfate) — As (10 mM sulfate)

What causes differences in patterns of Iron and Arsenic release caused by SRBs and FeRBs?

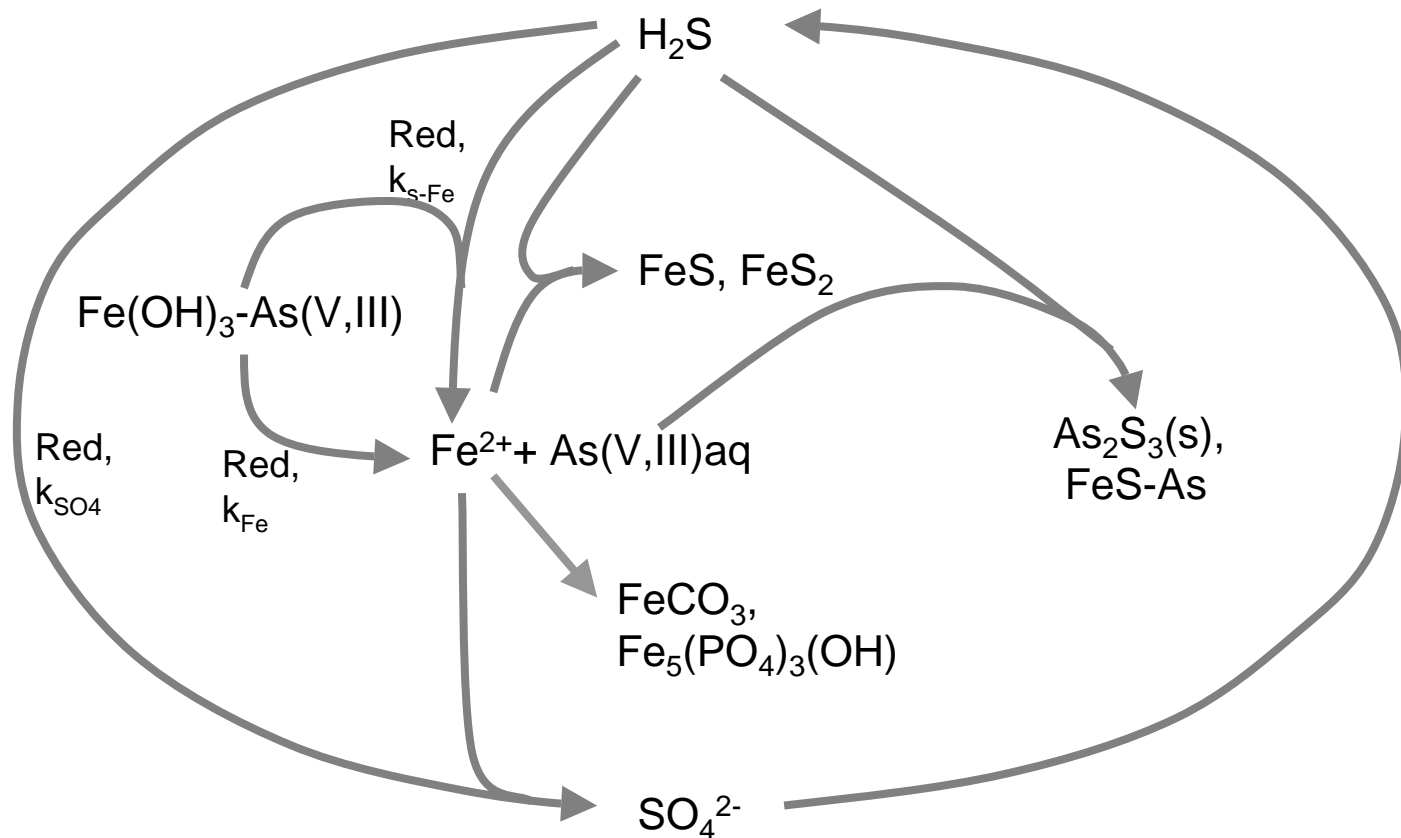


From Hansel, Fendorf, and Bostick 2004, unpublished

How do changes in $\delta^{34}\text{S}$ reflect sulfur cycling?



A more complete description of As fate



- It is necessary to include sulfate reduction to adequately describe arsenic concentrations.
- Kinetic processes are critical to regulating arsenic levels

Arsenic-Iron-Sulfur Cycling in 3 Field Sites

A photograph of a group of children and an adult standing in a field of tall grass. The children are of various ages, some looking towards the camera. The adult is holding a clipboard. In the background, there are several tall palm trees under a cloudy sky.

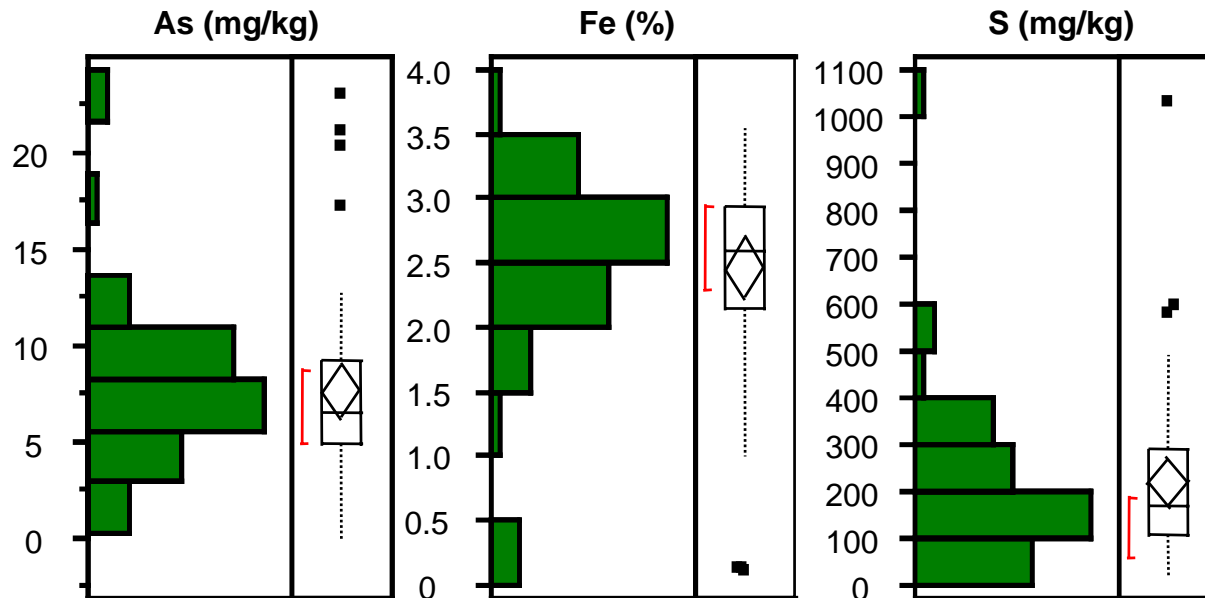
Coakley Superfund Site (NH)
Coeur d'Alene Mining District (ID)
Cambodian Groundwater Systems

Collaborators:

**Mickey Sampson (Resources
Development International,
Cambodia)**

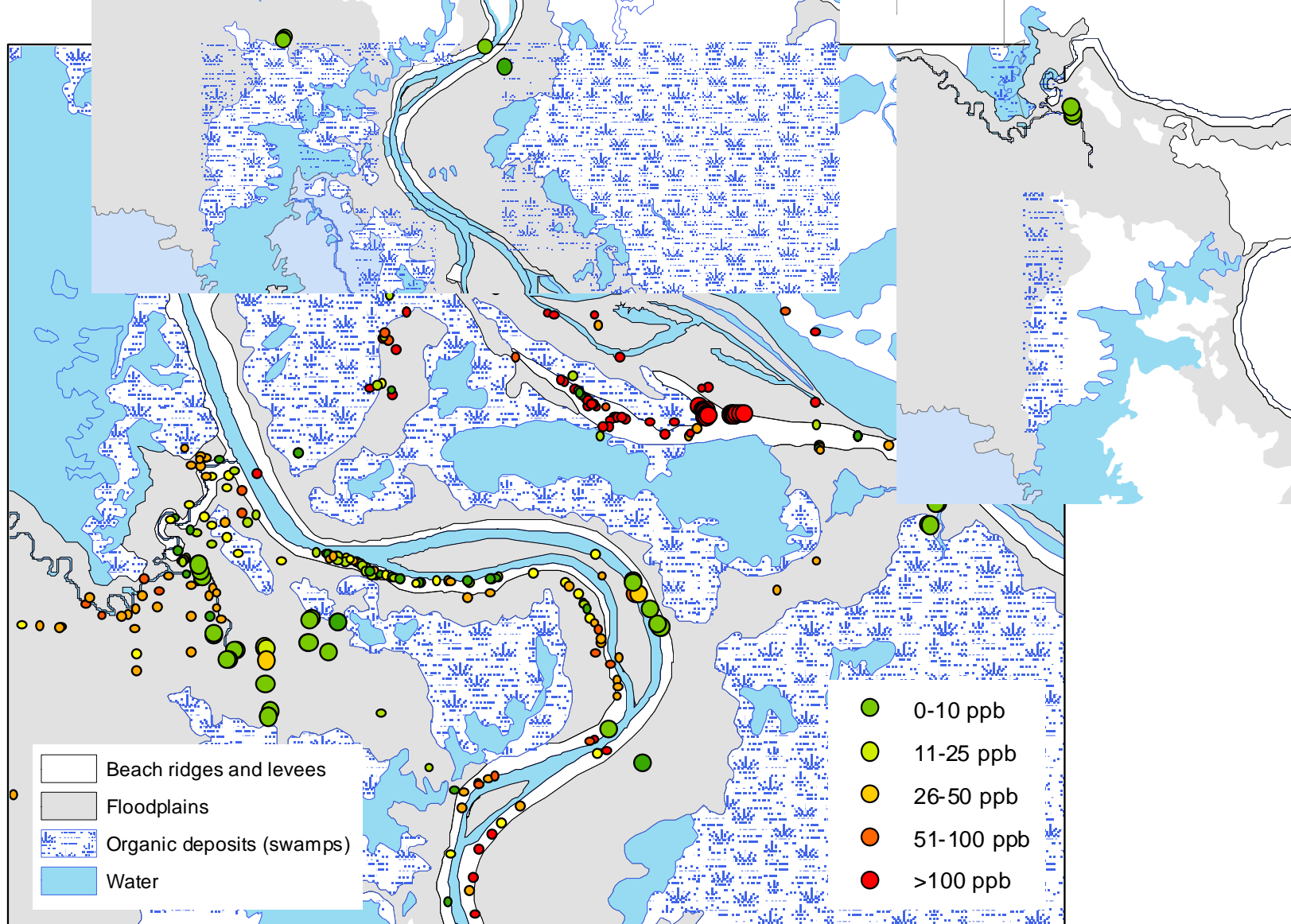
**Elizabeth Hadzima
Gretchen Gehrke
Nick Papacostas
Joshua Landis
Jamie de Lemos**

Soils and Sediments

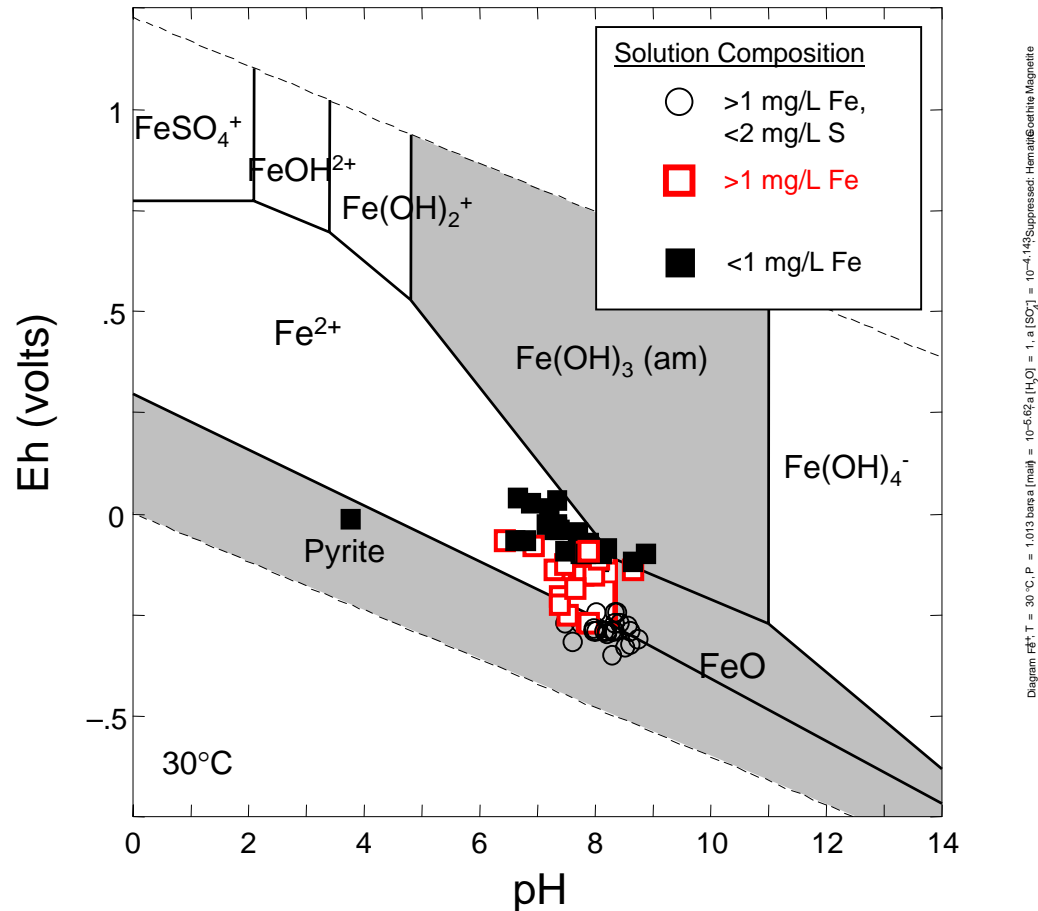


- Typical As, S and Fe levels
- Hard to determine composition of aquifer materials based on surficial environment

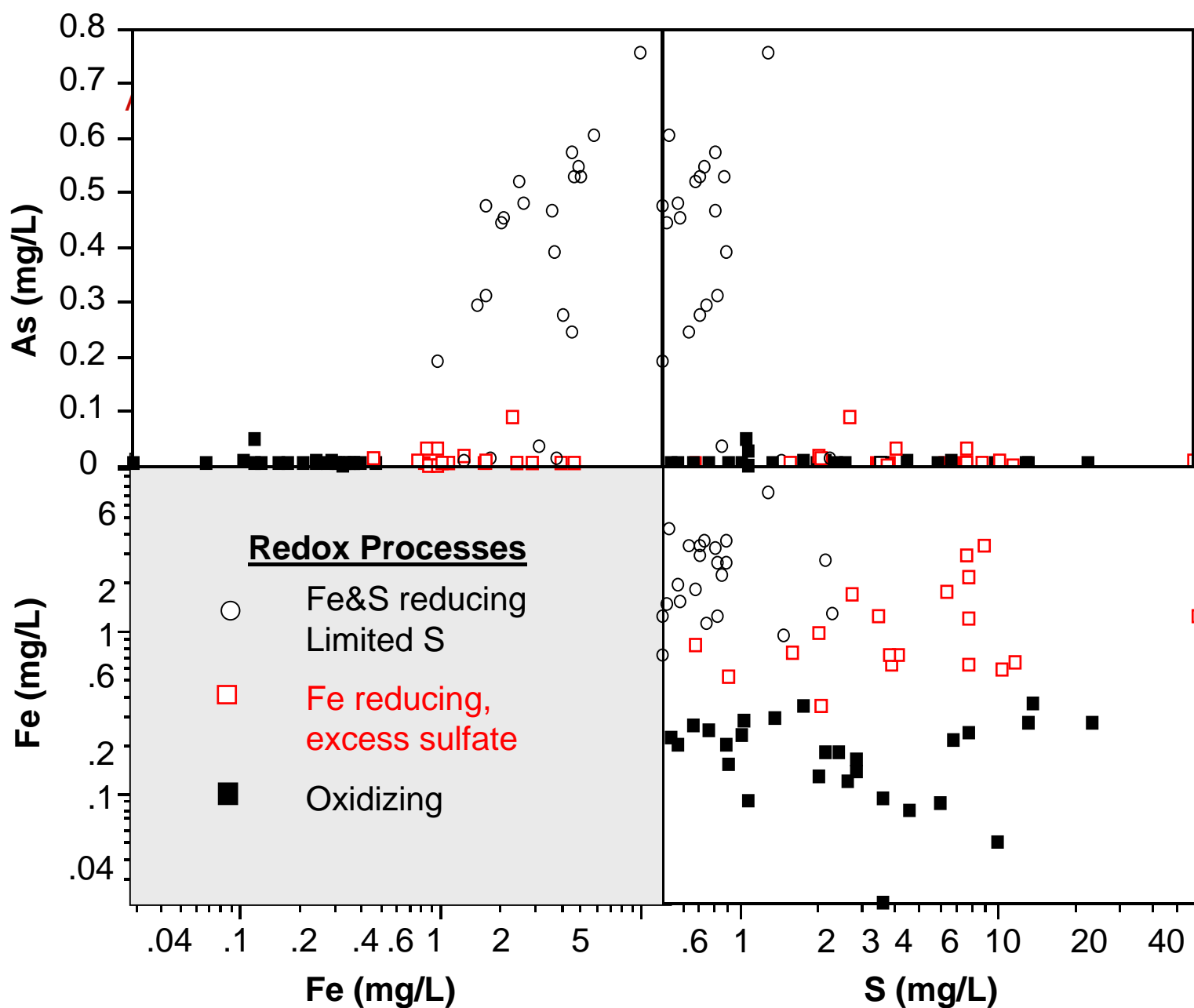
Arsenic and Regional Surficial Geology



Redox Processes: Sulfate and Iron reduction



- Redox Conditions indicate that sulfate reduction and/or Fe reduction is thermodynamically viable, concentration information indicates the extent to which they have occurred.



- Highly elevated As levels are most notably associated with waters high in Fe and low in sulfate

